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Remarks:
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(54) High fidelity Not1 restriction endonucleases

(57) *Inter alia*, a composition characterized in that it comprises: a restriction endonuclease enzyme having at least one artificially introduced mutation and an overall fidelity index (FI) improvement factor of at least 2, the restriction endonuclease being capable of cleaving a substrate with at least a similar cleavage activity to that of the restriction endonuclease absent the artificially introduced mutation, in a predetermined buffer, wherein the artificially introduced mutation is the product of at least one of a targeted mutation, saturation mutagenesis, or a mutation introduced through a PCR amplification procedure, and wherein the restriction endonuclease absent the artificially introduced mutation is NotI and the artificially introduced mutation is selected from: K176A; R177A; R253A; and K150A is disclosed.

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Description

BACKGROUND

[0001] Restriction endonucleases are enzymes that cleave doublestranded DNAs in a sequence-specific manner (Roberts, R.J., Proc Natl Acad Sci U S A, 102:5905-5908 (2005); Roberts, et al., Nucleic Acids Res, 31:1805-1812 (2003); Roberts, et al., Nucleic Acids Res, 33:D230-232 (2005); Alves, et al., Restriction Endonucleases, "Protein Engineering of Restriction Enzymes," ed. Pingoud, Springer-Verlag Berlin Heidelberg, New York, 393-407 (2004)). They are ubiquitously present among prokaryotic organisms (Raleigh, et al., Bacterial Genomes Physical Structure and Analysis, Ch.8, eds. De Bruijin, et al., Chapman & Hall, New York, 78-92 (1998)), in which they form part of restriction-modification systems, which mainly consist of an endonuclease and a methyltransferase. The cognate methyltransferase methylates the same specific sequence that its paired endonuclease recognizes and renders the modified DNA resistant to cleavage by the endonuclease so that the host DNA can be properly protected. However, when there is an invasion of foreign DNA, in particular bacteriophage DNA, the foreign DNA will be degraded before it can be completely methylated. The major biological function of the restriction-modification system is to protect the host from bacteriophage infection (Arber, Science, 205:361-365 (1979)). Other functions have also been suggested, such as involvement in recombination and transposition (Carlson, et al., Mol Microbiol, 27:671-676 (1998); Heitman, Genet Eng (N Y), 15:57-108 (1993); McKane, et al., Genetics, 139:35-43 (1995)).

[0002] The specificity of the approximately 3,000 known restriction endonucleases for their greater than 250 different target sequences could be considered their most interesting characteristic. After the discovery of the sequence-specific nature of the first restriction endonuclease (Danna, et al., Proc Natl Acad Sci U S A, 68: 2913-2917 (1971); Kelly, et al., J Mol Biol, 51:393-409 (1970)), it did not take long for scientists to find that certain restriction endonucleases cleave sequences which are similar but not identical to their defined recognition sequences under non-optimal conditions (Polisky, et al., Proc Natl Acad Sci U S A, 72:3310-3314 (1975); Nasri, et al., Nucleic Acids Res, 14:811-821 (1986)). This relaxed specificity is referred to as star activity of the restriction endonuclease. It has been suggested that water-mediated interactions between the restriction endonuclease and DNA are the key differences between specific complexes and star complexes (Robinson, et al., J Mol Biol, 234:302-306 (1993); Robinson, et al., Proc Natl Acad Sci U S A, 92: 3444-3448 (1995), Sidorova, et al., Biophys J, 87:2564-2576 (2004)).

[0003] Star activity is a problem in molecular biology reactions. Star activity introduces undesirable cuts in a cloning vector or other DNA. In cases such as forensic applications, where a certain DNA substrate needs to be cleaved by a restriction endonuclease to generate a unique fingerprint, star activity will alter a cleavage pattern profile, thereby complicating analysis. Avoiding star activity is also critical in applications such as strand displacement amplification (Walker, et al., Proc Natl Acad Sci U S A, 89:392-396 (1992)) and serial analysis of gene expression (Velculescu, et al., Science, 270:484-487 (1995)).

SUMMARY

[0004] In an embodiment of the invention, a composition is provided that includes a restriction endonuclease having at least one artificially introduced mutation and an overall fidelity index (FI) improvement factor of at least two, the restriction endonuclease being capable of cleaving a substrate with at least a similar cleavage activity to that of the restriction endonuclease absent the artificially introduced mutation in a predetermined buffer, the artificially introduced mutation being the product of at least one of a targeted mutation, saturation mutagenesis, or a mutation introduced through a PCR amplification procedure.

[0005] In a further embodiment of the invention, at least one of the artificially introduced mutations is a targeted mutation resulting from replacement of a naturally occurring residue with an oppositely charged residue. An Alanine or a Phenylalanine may replace the naturally occurring residue at the target site.

[0006] In a further embodiment of the invention, a composition of the type described above includes a restriction enzyme absent the artificially introduced mutation selected from the group consisting of: BamHI, EcoRI, ScaI, SalI, SphI, PstI, NcoI, NheI, SspI, NotI, SacI, PvuII, MfeI, HindIII, SbfI, EagI, EcoRV, AvrII, BstXI, PciI, HpaI, AgeI, BsmBI, BspQI, SapI, KpnI and BsaI.

[0007] Further embodiments of the invention include compositions listed in Table 4.

[0008] In a further embodiment of the invention, a DNA encoding any of the enzymes listed in Table 4 is provided, a vector comprising the DNA and a host cell for expressing the protein from the vector.

[0009] In an embodiment of the invention, a method is provided having the steps of (a) identifying which amino acid residues in an amino acid sequence of a restriction endonuclease having star activity are charged amino acids; (b) mutating one or more codons encoding one or more of the charged residues in a gene sequence encoding the restriction endonuclease; (c) generating a library of gene sequences having one or more different codon mutations in different charged residues; (d) obtaining a set of proteins expressed by the mutated gene sequences; and (e) determining an FI

in a predetermined buffer and a cleavage activity for each expressed protein.

[0010] An embodiment of the method includes the step of determining an overall FI improvement factor for proteins belonging to the set of proteins in a defined set of buffers where for example, the set of buffers contains NEB1, NEB2, NEB3 and NEB4 buffers.

[0011] An embodiment of the method includes the steps described above and additionally mutating codons encoding hydroxylated amino acids or amide amino acids in a same or subsequent step to that of mutating codons for the charged amino acids.

[0012] In an embodiment of the invention described above, the codons are mutated to an Alanine except for Tyrosine which is mutated to a Phenylalanine.

[0013] In a further embodiment, the overall FI improvement factor is improved using saturation mutagenesis of one or more of the mutated codon.

BRIEF DESCRIPTION OF THE DRAWINGS

[0014] For Figures 1-32:

[0015] The * symbol indicates the lane to its left that contains the lowest concentration of enzyme for which star activity is observed.

[0016] The # symbol refers to the lane showing incomplete cleavage, which is adjacent to and to the right side of the lane containing a concentration of enzyme sufficient for complete cleavage of the substrate.

[0017] The gray triangle denotes the serial decrease of restriction endonuclease concentration.

[0018] "U" denotes units of enzyme.

[0019] In each of the reactions described in Figures 1-32, the reaction mixture contains a volume of 3 μl unless otherwise specified of a buffer from New England Biolabs, Inc. (NEB), Ipswich, MA, (see Table 1 and NEB catalog), 3 μl unless otherwise specified of a specified restriction endonuclease in a diluent from NEB, Ipswich, MA (See Table 1 and NEB catalog) as well as variable volumes of specified substrate (containing 0.6 μg) substrate and a volume of water to bring the reaction mixture to a total of 30 μl. Reactions were conducted at 37°C for an incubation time of 1 hour. The results are analyzed on a 0.8% agarose gel. Where the overall volume of the reaction mix, amount of substrate, temperature of the reaction or incubation time varies from above, values are provided in the description of the figures.

[0020] The theoretical digestion pattern is provided on the right side of the gel for Figures 1, 5, 8, 11-18 and 20-32. Those substrates with only one restriction endonuclease site should be digested into one linear band from supercoiled form.

Figure 1 shows the determination of the FI for wild type (WT) ScaI by digesting 1.2 μl lambda DNA substrate (0.6 μg) with a two-fold serial dilution using diluent A of a preparation of WT ScaI (1,200 U) in NEB3 buffer and examining the digestion products on an agarose gel. The highest concentration of a restriction endonuclease with no star activity is shown with a solid arrow; and the minimum concentration giving rise to complete digestion of substrate is shown with a hollow arrow.

Figures 2A-D show the results of digesting 0.5 μl pUC19 substrate (0.5 μg) with WT BamHI or BamHI(E86P) enzyme in a three-fold serial dilution using diluent A for 1 hour at a starting concentration of 172 U or 512 U. The middle lane is the NEB 1 kb marker (New England Biolabs, Inc. (NEB), Ipswich, MA).

Fig. 2A shows results using NEB1 buffer.

Fig. 2B shows results using NEB2 buffer.

Fig. 2C shows results using NEB3 buffer.

Fig. 2D shows results using NEB4 buffer.

Figures 3A-B show a comparison of BamHI(E86P) activity over two time periods using 0.6 μl pBR322 substrate (which contains only 1 BamHI cleavage site) in NEB2 buffer using an initial concentration of 600 U of enzyme in a 2-fold serial dilution using diluent A.

Fig. 3A shows results in 1 hour.

Fig. 3B shows results in 14 hours.

Figures 4A-B show the cleavage of 0.6 μl pBR322 substrate in a 2-fold serial dilution of BamHI-HF (E163A/E167T) using diluent A after 14 hours incubation in two different buffers on an agarose gel.

Fig. 4A shows the results with NEB2 buffer with an initial concentration of 600 U of enzyme.

Fig. 4B shows the results with NEB1 buffer with an initial concentration of 2,400 U of enzyme.

Figures 5A-B show a comparison of cleavage reactions using BamHI-HF and WT BamHI in NEB4 buffer. The

reaction was carried out in NEB4 buffer using 1.2 μl lambda DNA substrate in a 2-fold serial dilution using diluent A.
Fig. 5A shows WT BamHI with a starting concentration of 1,200 U where the FI equals 4.
Fig. 5B shows BamHI-HF with a starting concentration of 2,400 U where the FI≥4000.

Figures 6A-D show a comparison of WT EcoRI and EcoRI(K62A) in NEB1-4 buffers in a 3-fold serial dilution using NEB diluent C. The reaction mixture contained 2 μl lambda DNA substrate (1 μg) in NEB1-4 buffers.
Fig. 6A shows the cleavage results following 2-fold serial dilution, 120 U WT EcoRI and 240 U of EcoRI (K62A) in NEB2 buffer.
Fig. 6B shows the cleavage results following 2-fold serial dilution, 120 U WT EcoRI and 240 U of EcoRI (K62A) in NEB4 buffer.
Fig. 6C shows the cleavage results following 2-fold serial dilution, 60 U WT EcoRI and 120 U of EcoRI (K62A) in NEB1 buffer.
Fig. 6D shows the cleavage results following 2-fold serial dilution, 120 U WT EcoRI and 60 U of EcoRI (K62A) in NEB3 buffer.

Figures 7A-C show the cleavage results with 2 different EcoRI mutants and WT EcoRI. The digestion of 100,000 U of enzyme and a 10-fold serial dilution thereof in diluent C over 10 hours using 0.6 μl of Litmus28 substrate in various buffers is shown. There is only one EcoRI cleavage site in Litmus28 substrate.
Fig. 7A: EcoRI mutant K62E in NEB4 buffer.
Fig 7B: EcoRI mutant K62A in NEB4 buffer.
Fig 7C: WT EcoRI in EcoRI buffer (see NEB catalog 2007-8).

Figures 8A-B shows a comparison of EcoRI-HF and WT EcoRI in NEB4 buffer. The reaction utilized 1.2 μl lambda DNA substrate in a 2-fold serial dilution using diluent C.
Fig. 8A: WT EcoRI with a starting concentration of 19,200 U reveals a FI=4 in NEB4 buffer.
Fig. 8B: EcoRI-HF with a starting concentration of 38,400 U reveals a FI=16,000 in NEB4 buffer.

Figures 9A-B shows a comparison of serial dilutions of WT ScaI (4.8 U), ScaI(H193A) (9.6 U), ScaI(S201F) (19.2 U), and ScaI(H193A/S201F) (19.2 U). Each sample was initially diluted by 1/10, followed by a 2-fold serial dilution in NEB2 buffer with the specified percentage of glycerol. Each reaction mixture contains 2 μl of lambda DNA substrate (1 μg).
Fig. 9A: 5% glycerol.
Fig 9B: 37% glycerol.

Figures 10A-D shows a comparison of WT ScaI and ScaI-HF (H193A/5201F). The enzymes (unit concentration as specified) were each diluted in a 2.5-fold serial dilution with diluent A. The reaction mixture contains 2 μl lambda DNA substrate and NEB1-4 buffers.
Fig. 10A: cleavage in NEB2 buffer.
Fig. 10B: cleavage in NEB4 buffer.
Fig. 10C: cleavage in NEB1 buffer.
Fig. 10D: cleavage in NEB3 buffer.

Figures 11A-H show the FI determination for SalI-HF and WT SalI. Both enzymes were diluted in 2-fold serial dilutions using diluent A. The reaction mixture contains 2μl HindIII-digested lambda DNA substrate.
Figs. 11A, B, C and D show a serial dilution of 1,200 U, 1,200 U, 300 U and 1,200 U of SalI-HF demonstrating a FI ≥1,000, FI ≥2,000, FI ≥500 and FI ≥2,000 in NEB1, 2, 3 and 4 buffers, respectively.
Figs. 11E, F, G, and H show a serial dilution of 19.2 U, 150 U, 9,600 U and 38.4 U of WT SalI demonstrating a FI =8, FI=1, FI=32 and FI=1 in NEB1, 2, 3 and 4 buffers, respectively.

Figures 12A-B show the results of a 2-fold serial dilution of SphI-HF (19,200 U) in diluent A and WT SphI (143,600 U) in diluent B reacted in NEB 4 buffer with 1.2 μl lambda DNA substrate.
Fig. 12A shows cleavage by WT SphI.
Fig. 12B shows cleavage by SphI-HF.

Figures 13A-B show cleavage of 1.2 μl lambda DNA substrate using 2-fold serial dilutions of PstI-HF (300 U and 150 U) and 2-fold serial dilutions of WT PstI (2,400 U and 4,800 U) in NEB3 and NEB4 buffers, respectively. Serial dilutions were performed in diluent C.
Fig. 13A shows cleavage by PstI-HF in NEB4 buffer (upper panel) and NEB3 buffer (lower panel).

Fig. 13B shows cleavage by WT PstI in NEB4 buffer (upper panel) and NEB3 buffer (lower panel).

Figures 14A-B show cleavage of 1.2 μl lambda DNA substrate using 2-fold serial dilutions of NcoI-HF (4,800 U and 600 U) and 2-fold serial dilutions of WT NcoI (4,800 U and 1,200 U) in NEB3 and NEB4 buffers, respectively. Serial dilutions were performed in diluent A.
Fig. 14A shows cleavage by NcoI-HF in NEB4 buffer (upper panel) and NEB3 buffer (lower panel).
Fig. 14B shows cleavage by WT NcoI in NEB4 buffer (upper panel) and NEB3 buffer (lower panel).

Figures 15A-B show cleavage of 1.2 μl pXba DNA substrate using 2-fold serial dilutions of NheI-HF (287,200 U and 76.8 U) and 2-fold serial dilutions of WT NheI (9,600 U and 300 U) in NEB3 and NEB4 buffers, respectively. Serial dilutions were performed in diluent A.
Fig. 15A shows cleavage by NheI-HF in NEB4 buffer (upper panel) and NEB3 buffer (lower panel).
Fig. 15B shows cleavage by WT NheI in NEB4 buffer (upper panel) and NEB3 buffer (lower panel.)

Figures 16A-B show cleavage of 1.2 μl lambda DNA substrate using 2-fold serial dilutions of SspI-HF (9,600 U and 38.4 U) and 2-fold serial dilutions of WT SspI (19,200 U and 19,200 U) in NEB3 and NEB4 buffers, respectively. Serial dilutions were performed in diluent C.
Fig. 16A shows cleavage by SspI-HF in NEB4 buffer (upper panel) and NEB3 buffer (lower panel).
Fig. 16B shows cleavage by WT SspI in NEB4 buffer (upper panel) and NEB3 buffer (lower panel).

Figures 17A-B show cleavage of 1.2 μl pXba DNA substrate using 2-fold serial dilutions of NotI-HF (287,200 U and 19,200 U) and 2-fold serial dilutions of WT NotI (19,200 U and 76,800 U) in NEB3 and NEB4 buffers, respectively. Serial dilutions were performed in diluent C.
Fig. 17A shows cleavage by NotI-HF in NEB4 buffer (upper panel) and NEB3 buffer (lower panel).
Fig. 17B shows cleavage by WT NotI I in NEB4 buffer (upper panel) and NEB3 buffer (lower panel).

Figures 18A-B show cleavage of 1.2 μl pXba DNA substrate using 2-fold serial dilutions of SacI-HF (4,800 U and 76.8 U) and 2-fold serial dilutions of WT SacI (19,200 U and 1200 U) in NEB3 and NEB4 buffers, respectively. Serial dilutions were performed in diluent A.
Fig. 18A shows cleavage by SacI-HF in NEB4 buffer (upper panel) and NEB3 buffer (lower panel).
Fig. 18B shows cleavage by WT SacI in NEB4 buffer (upper panel) and NEB3 buffer (lower panel).

Figures 19A-B show cleavage of 0.6 μl pBR322 DNA substrate using 2-fold serial dilutions of PvuII-HF (9,600 U and 19.2 U) and 2-fold serial dilutions of WT PvuII (19,200 U and 300 U) in NEB3 and NEB4 buffers, respectively. Serial dilutions were performed in diluent A.
Fig. 19A shows cleavage by PvuII-HF in NEB4 buffer (upper panel) and NEB3 buffer (lower panel)
Fig. 19B shows cleavage by WT PvuII in NEB4 buffer (upper panel) and NEB3 buffer (lower panel).

Figures 20A-B show cleavage of 1.2 μl lambda DNA substrate using 2-fold serial dilutions of MfeI-HF (300 U and 19.2 U) and 2-fold serial dilutions of WT MfeI (2,400 U and 38.4 U) in NEB3 and NEB4 buffers, respectively. Serial dilutions were performed in diluent A.
Fig. 20A shows cleavage by MfeI-HF in NEB4 buffer (upper panel) and NEB3 buffer (lower panel).
Fig. 20B shows cleavage by WT MfeI in NEB4 buffer (upper panel) and NEB3 buffer (lower panel).

Figures 21A-B show cleavage of 1.2 μl lambda DNA substrate using a 2-fold serial dilution of HindIII-HF (4,800 U and 1,200 U) and a 2-fold serial dilution of WT HindIII (9,600 U and 4,800 U) in NEB3 and NEB4 buffers, respectively. Serial dilutions were performed in diluent A.
Fig. 21A shows cleavage by HindIII-HF in NEB4 buffer (upper panel) and NEB3 buffer (lower panel).
Fig. 21B shows cleavage by WT HindIII in NEB4 buffer (upper panel) and NEB3 buffer (lower panel)

Figures 22A-B show cleavage of 1.2 μl lambda DNA substrate in a 2-fold serial dilution using diluent C of SbfI-HF (starting concentration: 76,800 U) and WT SbfI (starting concentration: 76,800 U) in NEB4 buffer.
Fig. 22A shows cleavage by WT SbfI.
Fig. 22B shows cleavage by SbfI-HF.

Figures 23A-B shows cleavage of 1.2 μl pXba DNA substrate in a 2-fold serial dilution of EagI-HF (1,200 U and 600 U) and a 2-fold serial dilution of WT EagI (150 U and 38.4 U) in NEB2 and NEB1 buffers, respectively, using diluent C.
Fig. 23A shows cleavage by EagI-HF in NEB2 buffer (upper panel) and NEB1 buffer (lower panel).

Fig. 23B shows cleavage by WT EagI in NEB2 buffer (upper panel) and NEB1 buffer (lower panel).

Figures 24A-B show cleavage of 1.2 μl pXba DNA substrate in a two-fold serial dilution using diluent A of EcoRV-HF (starting concentration: 38,400 U) and WT EcoRV (starting concentration: 2400 U) in NEB4 buffer.
Fig. 24A shows cleavage by WT EcoRV.
Fig. 24B shows cleavage by EcoRV-HF.

Figures 25A-B show cleavage of 1.2 μl T7 DNA substrate in a two-fold serial dilution using diluent A of AvrII-HF (starting concentration: 1,200 U) and WT AvrII (starting concentration: 1,200 U) in NEB4 buffer.
Fig. 25A shows cleavage by WT AvrII.
Fig. 25B shows cleavage by AvrII-HF.

Figures 26A-B show cleavage of 1.2 μl lambda DNA substrate by a two-fold serial dilution in diluent A of BstXI-HF (starting concentration: 300 U) and WT BstXI (starting concentration: 38.4 U) in NEB4 buffer. The reaction was performed at 55°C.
Fig. 26A shows cleavage by WT BstXI.
Fig. 26B shows cleavage by BstXI-HF.

Figures 27A-B show cleavage of 1.2 μl pXba DNA substrate in a two-fold serial dilution using diluent A of PciI-HF (starting concentration: 600 U) and WT PciI (starting concentration 300 U) in NEB4 buffer.
Fig. 27A shows cleavage by WT PciI.
Fig. 27B shows cleavage by PciI-HF.

Figures 28A-B show cleavage of 1.2 μl lambda DNA substrate in a two-fold serial dilution using diluent A of HpaI-HF (starting concentration: 4,800 U) and WT HpaI (starting concentration 9,600 U) in NEB2 buffer.
Fig. 28A shows cleavage by WT HpaI.
Fig. 28B shows cleavage by HpaI-HF.

Figures 29A-B show cleavage of 1.2 μl pXba DNA substrate in a two-fold serial dilution of AgeI-HF using diluent C (starting concentration: 600 U) and WT AgeI (starting concentration 600 U) in NEB4 buffer.
Fig. 29A shows cleavage by WT AgeI.
Fig. 29B shows cleavage by AgeI-HF.

Figures 30A-B show cleavage of 1.2 μl lambda DNA substrate in a two-fold serial dilution using diluent A of BsmBI-HF (starting concentration: 300 U) and WT BsmBI (starting concentration 4,800 U) in NEB4 buffer. The reaction is at 55°C.
Fig. 30A shows cleavage by WT BsmBI.
Fig. 30B shows cleavage by BsmBI-HF.

Figures 31A-B show the DNA sequence (SEQ ID NO:1) and protein sequence (SEQ ID NO:2) of MluCIM, respectively, for expression of EcoRI and MfeI.

Figure 32 shows the DNA sequence (SEQ ID NO:3) of Hpy166IIM for expression of SalI.

Figures 33A-B show the DNA sequence (SEQ ID NO:4) and protein sequence (SEQ ID NO:5) of MfeI, respectively.

Figures 34A-B show the DNA sequence (SEQ ID NO:6) and protein sequence (SEQ ID NO:7) of BstXI, respectively.

Figures 35A-B show the DNA sequence (SEQ ID NO:8) and protein sequence (SEQ ID NO:9 of M.BstXI, respectively.

Figures 36A-B show the DNA sequence (SEQ ID NO:10) and protein sequence (SEQ ID NO:11) of S.BstXI, respectively.

Figures 37A-B show the DNA sequence (SEQ ID NO:12) and protein sequence (SEQ ID NO:13) of PciI, respectively.

Figures 38A-B show the DNA sequence (SEQ ID NO:14) and protein sequence (SEQ ID NO:15) of M.PciI, respectively.

Figure 39 shows the DNA sequence (SEQ ID NO:17) encoding MI.EarI that recognizes CTCTTC and methylates at N4 cytosine or N6 adenine for cloning SapI and BspQI.

Figure 40 shows the DNA sequence (SEQ ID NO:18) encoding M2.EarI that recognizes CTCTTC and methylates at N4 cytosine or N6 adenine for cloning SapI and BspQI.

Figures 41A-B show agarose gels to determine the FI of 1 μl WT BspQI and 1 μl mutant (K279P/R388F) BspQI (starting concentrations 512 U and 1,024 U, respectively) by cleaving 1 μl pUC19 DNA substrate in a two-fold serial dilution using diluent A and NEB1 buffer plus 10% glycerol. The reaction was conducted at 50°C.

Figure 42 shows agarose gels to determine the FI of 5 μl WT SapI and 5 μl mutant (K273A) SapI (starting concentrations 32 U and 16 U, respectively) by cleaving pUC19 DNA substrate in a two-fold serial dilution using diluent A and NEB2 buffer plus 25% DMSO.

Figures 43A-B show catalytic and star activity of pXba by 5 μl WT KpnI and 5 μl D16N/E132A/D148E KpnI (initial concentrations 32 U and 256 U, respectively). The enzyme digested 2 μl pXba DNA substrate (0.5 μg) in a 2-fold serial dilution in NEB2 buffer using a diluent containing 10 mM Tris-HCl, pH 7.4, 50 mM KCl, 0.1 mM EDTA, 1 mM DTT and 50% glycerol with the total volume made up to 50 μl with water.

Fig. 45A shows the cleavage results of WT KpnI.

Fig. 45B shows the cleavage results of D16N/E132A/D148E KpnI.

Figure 44A-D show the amino acid sequences of the enzymes in Table 1 and the DNA sequences of the enzymes in Table 1 that have not been previously disclosed.

DETAILED DESCRIPTION OF THE EMBODIMENTS

[0021] Embodiments of the invention provide a general method for selecting for restriction endonucleases with desired characteristics. The general method relies on a suitable assay for determining whether the desired restriction endonuclease has been created. In particular an embodiment of the general method provides a systematic screening method with a set of steps. This method has been deduced by performing many hundreds of reactions using many restriction endonucleases. The majority of the examples provided herein relate to identifying restriction endonucleases with reduced star activity but with cleavage activity that is at least similar to the WT restriction endonuclease. However, it is expected that the same methodology can be applied successfully to modifying other properties of the restriction endonucleases relating, for example, to improved cleavage activity in desired buffers, thermostability, rate of reaction in defined conditions, etc.

[0022] As discussed above, an end point of interest is to transform restriction endonucleases with star activity into high fidelity restriction endonucleases with significantly reduced star activity. Star activity refers to promiscuity in cleavage specificity by individual restriction endonucleases. The terms "reduction in star activity" and "increase in fidelity" are used interchangeably here. Although restriction endonucleases are characterized by their property of cleaving DNA at specific sequences, some restriction endonucleases additionally cleave DNA inefficiently at secondary sites in the DNA. This secondary cleavage may occur consistently or may arise only under certain conditions such as any of: increased concentrations, certain buffers, temperature, substrate type, storage, and incubation time.

[0023] It is generally acknowledged that little is known about the complex environment generated by the hundreds of amino acids that constitute a protein and determine specificity. One approach in the prior art has been to utilize crystallography to identify contact points between an enzyme and its substrate. Nonetheless, crystallography has limitations with respect to freezing a structure in time in an unnatural chemical environment.

[0024] The rules that determine the contribution of amino acids at any site in the protein and the role played by the structure of the substrate molecule has proved elusive using existing analytical techniques. For example, it is shown here that mutating an amino acid in a restriction endonuclease can cause all or partial loss of activity.

[0025] In this context, no structural explanation has been put forward to explain why star activity could increase with high glycerol concentration (> 5% v/v), high enzyme to DNA ratio (usually > 100 units of enzyme per μg of DNA), low ionic strength (< 25 mM salt), high pH (> 8.0), presence of organic solvent (such as DMSO, ethanol), and substitution of Mg^{2+} with other divalent cations (Mn^{2+}, Co^{2+}). It was here recognized that because of the diversity of factors affecting star activity, it would be necessary to conduct comparisons of WT and mutant star activity under the same reaction conditions and in the same predetermined buffer and to develop a standard reaction condition in which any high fidelity enzyme must be capable of showing the described characteristics even if these characteristics were also observed in

other reaction conditions.

[0026] Present embodiments of the invention are directed to generating modified restriction endonucleases with specific improved properties, namely enhanced cleavage fidelity without significant reduction in overall cleavage activity or significant loss of yield from the host cells that make the protein. The methods that have been developed here for finding mutants with improved properties have resulted from exhaustive experimentation and the properties of the resultant enzymes have been defined in the context of specified conditions. The methods described herein may be used for altering the enzymatic properties of any restriction endonuclease under predetermined conditions, but are not limited to the specific defined conditions.

Restriction Endonuclease	Steps Used to Generate a High Fidelity Restriction Endonuclease
BamHI (Ex. 1)	Comparison of isoschizomer Targeted 22 residues to mutate to Ala. 14 mutants obtained, 3 had improved fidelity Saturation mutagenesis on 2 residues-K30 and E86 Recovered E86P as preferred mutant with greatest reduced star activity in selected buffers. Added mutations to E86P. Second round of mutation (Arg, Lys, His, Asp, Glu, Ser, Thr) to Ala and Tyr to Phe. Selected E167 and Y165 for saturation mutagenesis and selected E167T and Y165F. E163A/E167T was selected as preferred high fidelity mutant (BamHI-HF).
EcoRI (Ex. 2)	Comparison of isoschizomer Targeted 42 charged residues to mutate to Ala. No high fidelity mutants Second round of mutation: Target additional 32 charged residues to mutate to Ala: Identified K62A. Saturation mutagenesis on K62A. EcoRI(K62E) was selected as a preferred high fidelity mutant (EcoRI-HF).
ScaI (Ex. 3)	Comparison of isoschizomers. Targeted 58 charged residues to mutate to Ala. Identify 4 mutants Preferred mutant of 4 is (H193A/S201F). This is selected as a preferred high fidelity mutant (ScaI-HF)
SalI (Ex. 4)	Target 86 charged residues and mutate to Ala. SalI (R107A) was preferentially selected as a preferred high fidelity mutant (SalI-HF).
SphI (Ex. 5)	Target 71 charged residues and mutate to Ala. SphI (K100A) was preferentially selected as a preferred high fidelity mutant (SphI-HF)
PstI (Ex. 6)	Target 92 charged amino acids and mutate to Ala. PstI (D91A) was preferentially selected as a preferred high fidelity mutant (PstI-HF)
NcoI (Ex. 7)	Target 66 charged residues and mutate to Ala. NcoI (A2T /R31A) was preferentially selected as a preferred high fidelity mutant (NcoI-HF).
NheI (Ex. 8)	Target 92 charged residues and mutate to Ala. NheI (E77A) was preferentially selected as a preferred high fidelity mutant (NheI-HF)
SspI (Ex. 9)	Target 81 charged residues and mutate to Ala. No preferential mutants obtained. Target 95 residues to additional charged residues and hydroxylated residues to Ala except Tyr. Tyr mutated to Phe. SspI (Y98F) was preferentially selected as a preferred high fidelity mutant (SspI-HF)
NotI (Ex. 10)	Target 97 charged residues and mutate to Ala. K150A was preferentially selected as a preferred high fidelity mutant (NotI-HF)
SacI (Ex. 11)	Target 101 charged residues and mutate to Ala. SacI (Q117H/R200A) was preferentially selected as a preferred high fidelity mutant (SacI-HF) where Q117H was a carry over mutation from template with no affect on activity
PvuII (Ex. 12)	Target 47 charged residues and mutate to Ala. No preferred mutants obtained Target 19 hydroxylated residues - Ser/Thr and Tyr. Select T46A for further improvement

(continued)

Restriction Endonuclease	Steps Used to Generate a High Fidelity Restriction Endonuclease
	Saturation mutagenesis results in a preferred mutant T46G, T46H, T46K, T46Y. PvuII (T46G) was preferentially selected as a preferred high fidelity mutant (PvuII-HF)
MfeI (Ex. 13)	Target 60 charged residues and mutate to Ala. No preferred mutants obtained Target 26 hydroxylated residues and mutate to Ala except for Tyr which was changed to Phe. Target 38 residues (Cys, Phe, Met, Asn, Gln, Trp) and mutate to Ala Identify Mfe (Q13A/F35Y) as a preferred high fidelity mutant (MfeI-HF) where F35Y is carried from the template
HindIII (Ex. 14)	Target 88 charged residues and mutate to Ala. No preferred mutants obtained Target 103 residues (Cys Met Asn, Gln, Ser Thr Trp) and mutate to Ala and Tyr changed to Phe. Identify HindIII (K198A) as a preferred high fidelity mutant (HindIII-HF)
SbfI (Ex. 15)	Target 78 charged residues mutated to Ala Target 41 residues (Ser Thr) mutated to Ala /Tyr to Phe Target 55 residues of Cys, Phe, Met Asn, Gln, Trp to Ala SbfI (K251A) was selected as a preferred high fidelity mutant (SbfI-HF)
EagI (Ex. 16)	Target 152 residues (Asp, Glu, His, Lys, Arg, Ser, thr, Asn, and Gln changed to Ala and Tyr changed to Phe). EagI H43A was selected as a preferred high fidelity mutant (EagI-HF)
EcoRV (Ex. 17)	Target 162 residues (Cys,Asp, Glu, Phe, his, Lys, Met, Asn, Gln, Arg, Ser, Thr, to Ala and Trp to Phe) EcoRV (D19A/E27A) was selected as a preferred high fidelity mutant (EcoRV-HF)
AvrII (Ex. 18)	Target 210 residues (Cys,Asp, Glu, Phe, his, Lys, Met, Asn, Gln, Arg, Ser, Thr, to Ala and Trp to Phe) AvrII (Y104F) was selected as a preferred high fidelity mutant (AvrII-HF)
BstXI (Ex. 19)	Target 237 residues (Cys,Asp, Glu, Phe, his, Lys, Met, Asn, Gln, Arg, Ser, Thr, to Ala and Trp to Phe) BstXI (N65A) was selected as a preferred high fidelity mutant (BstXI-HF)
PciI (Ex. 20)	Target 151 residues (Cys,Asp, Glu, Phe, his, Lys, Met, Asn, Gln, Arg, Ser, Thr, to Ala and Trp to Phe) PciI (E78A/S133A) was selected as a preferred high fidelity mutant. (PciI-HF) This was spontaneous and not one of the 151 separate mutations
HpaI (Ex. 21)	Target 156 residues (Cys,Asp, Glu, Phe, his, Lys, Met, Asn, Gln, Arg, Ser, Thr, to Ala and Trp to Phe) HpaI (E56A) was selected as a preferred high fidelity mutant (HpaI-HF)
AgeI (Ex. 22)	Target 149 residues (Cys,Asp, Glu, Phe, his, Lys, Met, Asn, Gln, Arg, Ser, Thr, to Ala and Trp to Phe) AgeI (R139A) was selected as a preferred high fidelity mutant (AgeI-HF)
BsmBI (Ex. 23)	Target 358 residues (Cys,Asp, Glu, Phe, his, Lys, Met, Asn, Gln, Arg, Ser, Thr, to Ala and Trp to Phe) BsmBI(N185Y/R232A) was selected as a preferred high fidelity mutant (BsmBI (HF)
BspQI (Ex. 24)	Target 122 residues (Arg, Lys, His, Glu, Asp, Gln, Asn, Cys) Replace R at position 279 with Phe, Pro, Tyr, Glu, Asp or Leu. Preferred mutations were R388F and K279P. Created a double mutant BspQI(K279P/R388F) as preferred high fidelity mutant (BspQI-HF)
SapI	Find K273 and R380 in SapI corresponding to R388 and K279 in BspQI.

(continued)

Restriction Endonuclease	Steps Used to Generate a High Fidelity Restriction Endonuclease
(Ex. 25)	SapI (K273P/R380F) was selected as a preferred high fidelity mutant (SapI-HF)
KpnI (Ex. 26)	Target all residues (Asp, Glu, Arg, Lys, His, Ser, Thr, Tyr, Asn, Gln, Phe, Trp, Cys, Met) to Ala. More mutation was done on site D16 and D148. A combined D16N/E132A/D148E was selected as a preferred high fidelity mutant (KpnI-HF).
BsaI (Ex. 27)	Find 11 amino acids corresponding to the site in BsmBI. BsaI (Y231F) was selected as a preferred high fidelity mutant (BsaI-HF).

[0027] The method follows from the realization that amino acids responsible for cognate activity and star activity are different. The engineering of high fidelity restriction endonucleases described herein demonstrates that cognate activity and star activity can be separated and there are different critical amino acid residues that affect these different activities. The locations of amino acids that are here found to affect star activity are not necessarily found within the active site of the protein. The cleavage properties of any restriction endonuclease has been determined here for the first time by developing a criterion of success in the form of determining a FI (see also Wei et al. Nucleic Acid Res., 36, 9, e50 (2008)) and an overall fidelity index improvement factor.

[0028] An "overall fidelity index improvement factor" refers to the highest FI for a mutant with maximum cleavage activity divided by the highest FI of the corresponding WT endonuclease with maximum cleavage activity within a selected set of buffers. The selected set may be of any size greater than one but practically will contain less than 10 different buffers and more preferably contains 4 buffers. The set may also include less than 4 buffers. The overall FI improvement factor of at least two should preferably be applicable for any mutant restriction endonuclease in the claimed invention additionally but not exclusively to the set of buffers consisting of NEB1, NEB2, NEB3 and NEB4.

[0029] A "similar cleavage activity" can be measured by reacting the same amount of enzyme with the same amount and type of substrate under the same conditions and visually comparing the cleavage profiles on a gel after electrophoresis such that the amount of cleavage product appears to be the same within a standard margin of error and wherein the quantitative similarity is more than 10%.

[0030] "Artificial" refers to "man-made".

[0031] "Standard conditions" refers to an overall FI improvement factor calculated from results obtained in NEB1-4 buffers.

[0032] The general method described herein has been exemplified with 27 restriction endonucleases: AgeI, AvrII, BamHI, BsaI, BsmBI, BspQI, BstXI, EagI, EcoRI, EcoRV, HindIII, HpaI, KpnI, MfeI, NcoI, NheI, NotI, PciI, PstI, PvuII, SacI, SalI, SapI, SbfI, ScaI, SphI and SspI restriction endonucleases. However, as mentioned above, the method is expected to be effective for the engineering of any restriction endonuclease that has significant star activity.

[0033] Embodiments of the method utilize a general approach to create mutant restriction endonucleases with reduced star activity. For certain enzymes, it has proven useful to mutate charged residues that are determined to be conserved between two isoschizomers (see for example SapI in Example 25). In general, however, the method involves a first step of identifying all the charged and polar residues in a protein sequence for the endonuclease. For example, charged amino acids and polar residues include the acidic residues Glu and Asp, the basic residues His, Lys and Arg, the amide residues Asn and Gln, the aromatic residues Phe, Tyr and Trp and the nucleophilic residue Cys. Individual residues are targeted and mutated to an Ala and the products of these targeted mutations are screened for the desired properties of increased fidelity. If none of the mutants obtained provide a satisfactory result, the next step is to target mutations to all the hydroxylated amino acids, namely, Ser, Thr and Tyr, the preferred mutation being Ser and Thr to Ala and Tyr to Phe. It is also possible to target mutations to both classes of residues at one time as was done for Examples 16-23. The mutation to Ala may be substituted by mutations to Val, Leu or Ile.

[0034] After these analyses, if one or more of the preferred mutants generated in the above steps still have substandard performance under the selected tests, these mutants can be selected and mutated again to each of the additional possible 18 amino acids. This is called saturation mutagenesis. Saturation mutagenesis provided the preferred high fidelity mutants for EcoRI (Example 2), BamHI in part (Example 1) and PvuII (Example 12). Depending on the results of saturation mutagenesis, the next step would be to introduce additional mutations either targeted or random or both into the restriction endonuclease. In Example 11, SacI-HF includes a random mutation generated fortuitously during inverse PCR. In Example 20, PciI-HF resulted from a random mutation and not from targeted mutations. In Example 26, BspQI-HF contains two mutations that were found to act synergistically in enhancing fidelity.

[0035] The use of various methods of targeted mutagenesis such as inverse PCR may involve the introduction of non-

target mutations at secondary sites in the protein. These secondary mutations may fortuitously provide the desired properties (see Example 20). It is desirable to examine those mutated enzymes with multiple mutations to establish whether all the mutations are required for the observed effect. In Example 11, Q117H in the double mutant had no effect on activity. In Example 20, the additional spontaneous mutation appears to be solely responsible for the observed improved fidelity, whereas in Example 24, the individual mutations acted synergistically.

[0036] In some cases, a mutation may provide an additional advantage other than improved fidelity (see for example BamHI in which either E163A or P173A causes the enzyme to become more thermolabile).

[0037] The high fidelity/reduced star activity properties of the mutants provided in the Examples were selected according to their function in a set of standard buffers. Other mutations may be preferable if different buffer compositions were selected. However, the same methodology for finding mutants would apply. Table 4 lists mutations which apply to each restriction endonuclease and provide an overall FI improvement factor in the standard buffer.

[0038] The engineering of the high fidelity restriction endonucleases to provide an overall FI improvement factor of at least 2 involves one or more of the following steps:

1. Assessment of the star activity of the WT restriction endonuclease

[0039] In an embodiment of the invention, the extent of star activity of a restriction endonuclease is tested by means of the following protocol: the endonuclease activity is determined for an appropriate substrate using a high initial concentration of a stock endonuclease and serial dilutions thereof (for example, two-fold or three-fold dilutions). The initial concentration of restriction endonuclease is not important as long as it is sufficient to permit an observation of star activity in at least one concentration such that on dilution, the star activity is no longer detected.

[0040] An appropriate substrate contains nucleotide sequences that are cleaved by cognate endonuclease activity and where star activity can be observed. This substrate may be the vector containing the gene for the restriction endonuclease or a second DNA substrate. Examples of substrates used in Table 2 are pBC4, pXba, T7, lambda, and pBR322.

[0041] The concentration of stock restriction endonuclease is initially selected so that the star activity can be readily recognized and assayed in WT and mutated restriction endonucleases. Appropriate dilution buffers such as NEB diluent A, B or C is selected for performing the serial dilutions according to guidelines in the 2007-08 NEB catalog. The serially diluted restriction endonuclease is reacted with a predetermined concentration of the appropriate substrate in a total reaction volume that is determined by the size of the reaction vessel. For example, it is convenient to perform multiple reactions in microtiter plates where a 30 μl reaction mixture is an appropriate volume for each well. Hence, the examples generally utilize 0.6 μg of substrate in 30 μl, which is equivalent to 1 μg of substrate in 50 μl. The amount of substrate in the reaction mixture is not critical, but it is preferred that it be constant between reactions. The cleavage reaction occurs at a predetermined temperature (for example 25°C, 30°C, 37°C, 50 °C, 55°C or 65°C) for a standard time such as one hour. The cleavage products can be determined by any standard technique, for example, by 0.8% agarose gel electrophoresis to determine the fidelity indices as defined above.

[0042] Not all restriction endonucleases have significant star activity as determined from their FI. However, if an endonuclease has a highest FI of no more than about 250 and a lowest FI of less than 100, the restriction endonuclease is classified as having significant star activity. Such endonucleases are selected as a target of enzyme engineering to increase fidelity for a single substrate. In some cases, the restriction endonucleases with both FI over about 500 and FI less than about 100 are also engineered for better cleavage activity.

[0043] Table 2 below lists the FI of some engineered restriction endonucleases before engineering. All samples were analyzed on 0.8% agarose gel.

Table 2

Enzyme	Diluent (NEB)***	Substrate*	Temp °C	FI-1 **	FI-2 **	FI-3 **	FI-4 **
AgeI	C	pXba	37	16(1)	8(½)	64(1/8)	8(½)
AvrII	B	T7	37	64(1)	8(1)	32(¼)	32(1)
BamHI	A	λ	37	4(½)	4(1)	32(1)	4(½)
BsaI	B	pBC4	50	8(1/4)	120(1)	16(1/4)	32(1)
BsmBI	B	λ	55	1(1/8)	8(½)	120(1)	4(¼)
BspQI	B	λ	50	2(1/8)	16(1)	32(1)	4(½)
BstXI	B	λ	55	2(½)	2(½)	2(1/8)	4(1)
EagI	B	pXba	37	4(¼)	8(½)	250(1)	16(1)

(continued)

Enzyme	Diluent (NEB)***	Substrate*	Temp °C	FI-1 **	FI-2 **	FI-3 **	FI-4 **
EcoRI	C	λ	37	250(½)	4(1)	250(1)	4(1)
EcoRV	A	pXba	37	32(1/16)	120(½)	1000(1)	64(¼)
HindIII	B	λ	37	32(¼)	250(1)	4000(¼)	32(½)
HpaI	A	λ	37	32(1/16)	1(¼)	2(1/8)	16(1)
KpnI	A	pXba	37	16(1)	16(¼)	8(1/16)	4(½)
MfeI	A	λ	37	32(1)	16(1/8)	8(1/16)	32(1)
NcoI	A	λ	37	120(1)	32(1)	120(¼)	32(1)
NheI	C	pXba	37	32(1)	120(¼)	120(1/8)	32(1)
NotI	C	pXba	37	≥32000(1/1 6)	64(1)	500(1)	32(¼)
PciI	A	pXba	37	2000(½)	16(¼)	120(1)	8(1/8)
PstI	C	λ	37	64(1)	32(1)	120(1)	8(½)
PvuII	A	pBR322	37	250(1)	16(¼)	8(1/32)	¼(1)
SacI	A	pXba	37	120(1)	120(½)	120(1/32)	32(½)
SalI	A	λ (H3)	37	8(1/500)	1(1/16)	32(1)	1(1/120)
SapI	C	λ	37	16(¼)	64(½)	32(¼)	16(1)
SbfI	A	λ	37	32(1)	8(¼)	8(1/16)	8(½)
ScaI	A	λ	37	1/16(1/37)	1/8(1)	4(½)	1/64(1/16)
SphI	B	λ	37	64(1)	32(1)	64(¼)	16(½)
SspI	C	λ	37	64(1)	16(1)	32(¼)	16(1)

* Substrate: λ is lambda phage DNA; λ (H3) is HindIII-digested lambda phage DNA; pXba is pUC19 with XbaI-digested fragment of Adeno Virus; pBC4: a shorter version of pXba; T7: T7 DNA

**FI-1 to FI-4: fidelity index of the enzyme in NEBuffer 1, 2, 3 and 4. The number in parenthesis is a value for relative cleavage activity of the mutant restriction endonuclease in a specified buffer in a set of buffers compared with the "best" cleavage activity of the same mutant restriction endonuclease in any of the buffers in the set of buffers.

[0044] The compositions of NEB buffers follow:

NEB1: 10 mM Bis Tris Propane-HCl, 10 mM $MgCl_2$, 1 mM dithiothreitol (pH 7.0 at 25°C);
NEB2: 50 mM NaCl, 10 mM Tris-HCl, 10 mM $MgCl_2$, 1 mM dithiothreitol (pH 7.9 at 25°C);
NEB3: 100 mM NaCl, 50 mM Tris-HCl, 10 mM $MgCl_2$, 1mM dithiothreitol (pH7.9 at 25°C);
NEB4: 50 mM potassium acetate, 20 mM Tris-acetate, 10 mM magnesium acetate, 1 mM dithiothreitol (pH7.9 at 25°C).

[0045] *** The compositions of NEB diluents follow. (Using diluents in the dilution instead of water will keep the glycerol concentration in the reaction as a constant.)
Diluent A: 50 mM KCl, 10 mM Tris-HCl, 0.1 mM EDTA, 1 mM dithiothreitol, 200 mg/ml BSA. 50% glycerol (pH7.4 at 25°C);
Diluent B: 300 mM NaCl, 10 mM Tris-HCl, 0.1 mM EDTA, 1mM dithiothreitol, 500 mg/ml BSA, 50% glycerol (pH7.4 at 25°C);
Diluent C: 250 mM NaCl, 10 mM Tris-HCl, 0.1 mM EDTA, 1 mM dithiothreitol, 0.15% Triton X-100, 200 mg/ml BSA, 50% glycerol (pH 7.4 at 25°C).

2. Construction of high expression host cell strains

[0046] It is convenient if a host cell is capable of over-expressing the mutant restriction endonuclease for which reduced star activity is sought. If the restriction enzyme is highly expressed in E. coli, the star activity can be readily detected in

the crude extract, which simplifies the screening for the high fidelity restriction endonuclease. However, the mutated restriction endonuclease can be expressed in any host cell providing that the host cell is protected in some way from toxicity arising from enzyme cleavage. This might include: the presence of a methylase; production in a compartment of the cell which provides a barrier to access to the genome (such as an inclusion body or the periplasm); in vitro synthesis; production in an emulsion (see U.S. patent application serial no. 12/035,872) absence of cleavage sites in the host genome; manufacture of the enzyme in component parts subject to intein mediated ligation (see U.S. Patent No. 6,849,428), etc.

[0047] Over-expression of the mutated restriction endonucleases for purposes of production can be achieved using standard techniques of cloning, for example, use of an E. coli host, insertion of the endonuclease into a pUC19-derived expression vector, which is a high copy, and use of a relatively small plasmid that is capable of constant expression of recombinant protein. The vector may preferably contain a suitable promoter such as the lac promoter and a multicopy insertion site placed adjacent to the promoter. Alternatively, a promoter can be selected that requires IPTG induction of gene expression. If the activity in the crude extract is not sufficient, a column purification step for the restriction endonuclease in crude extract may be performed.

3. Mutagenesis of restriction endonuclease

[0048] DNA encoding each charged or polar group in the restriction endonuclease may be individually targeted and the mutated DNA cloned and prepared for testing. Multiple mutations may be introduced into individual restriction endonuclease genes. Targeted mutagenesis of restriction endonucleases may be achieved by any method known in the art. A convenient method used here is inverse PCR. In this approach, a pair of complementary primers that contains the targeted codon plus a plurality of nucleotides (for Example 18 nt) on both the 5' and 3' side of the codon is synthesized. The selection of suitable primers can be readily achieved by reviewing the gene sequence of the endonuclease of interest around the amino acid residue of interest. Access to gene sequences is provided through REBASE and GenBank. The sequences for the endonucleases described herein in the Examples are provided in Figures 31 to 38 and 44. The template for PCR is a plasmid containing the restriction endonuclease gene. The polymerase is preferably a high fidelity polymerase such as Vent® or Deep Vent™ DNA polymerase. By varying the annealing temperature and Mg^{2+} concentration, successful introduction of most mutations can be achieved. The PCR amplification product is then purified and preferably digested by DpnI. In an embodiment of the invention, the digested product was transformed into competent host cells (for example, *E. coli*), which have been pre-modified with a corresponding methylase. Colonies from each mutant were picked and grown under similar conditions to those in which the WT is grown (for example, using similar growth medium, drug selection, and temperature). The resulting restriction endonucleases were screened for reduced star activity.

4. Screening for mutant restriction endonucleases with reduced star activity

[0049] Conditions such as buffer composition, temperature and diluent should be defined for determining star activity in a mutant restriction endonuclease. Tables 2 and 3 show the FI of recombinant endonucleases before and after mutation in four different buffers using three different diluents at 37°C. Accordingly, it is possible to determine which mutants have an overall desirable improved fidelity index factor of at least 2, more than 10, at least 50 or more than 500 and to select enzymes as preferred high fidelity mutants.

[0050] In an embodiment of the invention, the mutant restriction endonucleases were screened for activity in normal buffer conditions (no more than 5% glycerol) first. For those mutants with at least about 10% of activity of WT restriction endonuclease, activity was also determined in star activity promotion conditions that promoted star activity, for example, high glycerol concentration and optionally high pH. Preferably, the mutant with the least star activity but with acceptable cognate activity in normal buffers is selected. Plasmid can then be extracted and sequenced for the confirmation of the mutant. In some cases, the star activity is not easily measured, even with high glycerol and high pH conditions. Instead, the activity in different buffers is measured and compared, and the one with the highest cleavage activity ratio in NEB4 compared with NEB3 can be tested further for star activity improvement.

5. Saturation mutagenesis on one single residue

[0051] As described in the previous section, the first step is to mutate a target amino acid in the restriction endonuclease to Ala. If the results are not satisfactory, saturation mutagenesis is performed. This is preferably performed by one of two methods. One method is to change the intended codon into NNN. After mutagenesis, multiple colonies are assayed under normal conditions and under conditions that promote star activity. Alternatively, a different codon can be selected for mutagenesis of each of the targeted amino acids for example: Ala: GCT; Cys: TGC; Asp: GAC; Glu: GAA; His: CAC; Ile: ATC; Lys: AAA; Leu: CTG; Met: ATG; Asn: AAC; Pro: CCG; Gln: CAG; Arg: CGT; Ser: TCC; Thr: ACC; Val: GTT;

Trp: TGG and Tyr: TAC

6. Combination

[0052] More than one mutation can be introduced into the restriction endonuclease gene if a single mutation does not sufficiently reduce the star activity. Mutation combination and saturation mutagenesis can be performed in any order.

7. Mutant purification and assessment of the improvement

[0053] The high fidelity mutants may be purified in a variety of ways including use of different chromatography columns. For normal quality assessment, one FPLC heparin column is enough to eliminate the DNA and non-specific nucleases from the preparation. Multiple columns including ion exchange, hydrophobic, size exclusion and affinity columns can be used for further purification.

[0054] Purified high fidelity restriction endonucleases are measured for FI in four NEB buffers and compared with the FIs of the WT restriction endonuclease. The ratio of FI for the high fidelity restriction endonuclease in its optimal buffer to that of WT is the overall improvement factor.

Table 3 - FI* for exemplified restriction endonucleases

Enzyme	Diluent (NEB)	Substrate *	Temp °C	FI-1 **	FI-2 **	FI-3 **	FI-4 **
AgeI-HF	C	Xba	37	≥500(1)	≥250(½)	≥16(1/16)	≥250(1)
AvrII-HF	B	T7	37	500(1)	≥500(½)	≥16(1/64)	≥1000(1)
BamHI-HF	A	λ	37	≥4000(1)	≥4000(1)	≥250(1/16)	≥4000(1)
BsaI	B	pBC4	50	≥4000(1/2)	≥8000(1)	120(1)	≥8000(1)
BsmBI	B	λ	55	2(1)	≥500(1)	≥64(1/8)	≥500(1)
BspQI-HF	A	pUC19	50	≥1000(1/4)	≥1000(1/4)	≥64(1/64)	≥4000(1)
BstXI-HF	A	λ	55	≥120(½)	≥250(1)	≥16(1/16)	≥250(1)
EagI-HF	C	pXba	37	250(½)	250(1)	250(½)	500(1)
EcoRI-HF	C	λ	37	2000(1/8)	4000(¼)	250(1/250)	16000(1)
EcoRV-HF	A	pXba	37	≥16000(¼)	>64000(1)	≥32000(½)	≥64000(1)
HindIII-HF	B	λ	37	≥16000(¼)	≥64000(1)	≥16000(¼)	≥32000(½)
HpaI-HF	A	λ	37	≥32(1/32)	≥2000(1)	2(1/8)	≥2000(½)
KpnI-HF	A	pXba	37	≥4000(1)	≥1000(1/4)	≥64(1/64)	≥4000(1)
MfeI-HF	A	λ	37	≥1000(1)	≥250(¼)	≥16(1/64)	≥500(½)
NcoI-HF	A	λ	37	≥4000(¼)	≥4000(¼)	≥1000(1/16)	≥64000(1)
NheI-HF	C	pXba	37	≥128000(1)	≥4000(1/32)	≥32(1/2000)	≥32000(½)
NotI-HF	C	pXba	37	≥8000 (1/16)	≥128000(1)	≥4000(1/64)	≥64000(½)
PciI-HF	A	pXba	37	NC	≥2000(1)	≥2000(1)	≥1000(1)
PstI-HF	C	λ	37	1000(1/8)	4000(½)	4000(¼)	4000(1)
PvuII-HF	A	pBR322	37	≥250 (1/120)	≥2000(1/16)	≥250(1/120)	500(1)
SacI-HF	A	pXba	37	≥32000(1)	≥16000(½)	≥500(1/64)	≥32000(1)
SalI-HF	A	λ (H3)	37	≥8000(1/8)	≥64000(1)	≥4000(1/16)	≥32000(½)
SbfI-HF	C	λ	37	1000(1)	120(½)	8(1/32)	250(1)
ScaI-HF	A	λ	37	4000(1/8)	1000(1)	2000(1/32)	1000(1)

(continued)

Enzyme	Diluent (NEB)	Substrate *	Temp °C	FI-1 **	FI-2 **	FI-3 **	FI-4 **
SphI-HF	B	λ	37	4000(1/8)	2000(1/16)	250(1/250)	8000(1)
SspI-HF	C	λ	37	≥4000(½)	120(½)	≥32(1/128)	500(1)

* The FI is a ratio of the highest concentration that does not show star activity to the lowest concentration that completes digestion of the substrate.

** The number in parenthesis is a value for relative cleavage activity of the mutant restriction endonuclease in a specified buffer in a set of buffers compared with the greatest cleavage activity of the same mutant restriction endonuclease in any of the buffers in the set of buffers.

Table 4 - Mutations providing restriction endonucleases with high fidelity

Restriction Endonuclease	Examples of mutants with overall improved FI factor≥2
AgeI	R139A; S201A*
AvrII	Y104F; M29A; E96A; K106A; S127A; F142A
BamHI	E163A/E167T; K30A; E86A; E86P; K87A; K87E; K87V; K87N; P144A; Y165F; E167A; E167R; E167K; E167L; E167I K30A/E86A; E86A/K106A; K30A/E86A/K106A; K30A/K87A; E86P/K87E; E86A/Y165F; K30A/E167A; E163S/E170T/P173A; E163S/E170T/P173A; E86P/K87T/K88N/ E163S/E170T/P173A; E86P/K87R/K88G/E163S/E170T/P173A;E86P/K87P/K88R/ E163S/E170T/P173A/E211K; E86P/K87T/K88R/ E163S/E170T/P173A/N158S; E86P/K87S/K88P/ E163S/E170T/P173A; E86P/K87G/K88S/ E163S/E170T/P173A; E86P/K87R/K88Q/ E163S/E170T/P173A; E86P/K87W/K88V; E86P/P173A
BsaI	Y231F
BsmBI	N185Y/R232A; H230A; D231A; R232A;
BspQI	K279P/R388F; K279A; K279F; K279P; K279Y; K279E; K279D R388A; R388F; R388Y; R388L; K279P/R388F; K279A/R388A; D244A
BstXI	N65A; Y57F; E75A; N76A; K199A;
EagI	H43A
EcoRI	K62A; K62S; K62L; R9A; K15A; R123A; K130A; R131A; R183A; S2Y; D135A; R187A; K62E
EcoRV	D19A; E27A; D19A/E27A
HindIII	S188P/E190A; K198A
HpaI	Y29F; E56A
KpnI	D148E; D16N/R119A/D148E; D2A/D16N/D148E; D16N/E134A/D148E; D16N/E132A/D148E
MfeI	Y173F; Q13A/F35Y
NcoI	D56A; H143A; E166A; R212A; D268A; A2T/R31A
NheI	E77A
NotI	K176A; R177A; R253A; K150A
PciI	E78A/S133A
PstI	E204G; K228A; K228A/A289V; D91A
PvuII	T46A; T46H; T46K; T46Y; T46G
SacI	Q117H/R154A/L284P; Q117H/R200A

(continued)

Restriction Endonuclease	Examples of mutants with overall improved FI factor≥2
SalI	R82A; K93A; K101A; R107A
SapI	K273P; R380A; K273P/R380A
SbfI	K251A
ScaI	R18A; R112A; E119A; H193A; S201F; H193A/S201F
SphI	D91A; D139A; D164A; K100A
SspI	H65A; K74A; E78A; E85A; E89A; K109A; E118A; R177A; K197A; Y98F
The mutations for each enzyme are separated by a semicolon.	

EXAMPLES

[0055] Where amino adds are referred to by a single letter code, this is intended to be standard nomenclature. The key to the code is provided for example in the NEB catalog 2007/2008 on page 280.

[0056] Plasmids used for cloning and as substrates have sequences as follows:

[0057] pLaczz2 (SEQ ID NO:102), pSyx20-laclq (SEQ ID NO:105), pBC4 (SEQ ID NO:103), pXba (SEQ ID. NO:104) and pAGR3 (SEQ ID NO:106). pACYC is described in GenBank XO 6403, T7 in GenBank NC001604, pUC18 in GenBank L09136, and pRRS in Skoglund et al. Gene, 88:1-5 (1990. pSX33 was constructed by inserting lacI gene into pLG339 at EcoRI site. pLG339 is described in Stoker, et al. Gene 19, 335-341 (1982).

[0058] All buffers identified as NEB buffers used herein are obtainable from New England Biolabs, Inc. (NEB), Ipswich, MA.

Example Engineering of high fidelity NotI

1. Expression of NotI

[0059] NotI has significant star activity in NEB4 buffer and less in NEB3 buffer. NotI was engineered to reduce star activity in any NEB buffer. NotI was expressed in competent *E. coli* transformed with pACYC184-EagIM and placzz2-NotIR. The cells were grown at 37°C for overnight in the LB with Amp and Cam.

2. Mutagenesis of NotI

[0060] All 97 charged residues in NotI were mutated to Ala as the following residues: 2, 4, 8, 10, 17, 21, 22, 26, 31, 34, 35, 36, 49, 52, 57, 59, 62, 72, 74, 75, 77, 84, 87, 96, 97, 105, 117, 121, 122, 125, 126, 129, 130, 133, 140, 141, 145, 150, 152, 156, 160, 165, 167, 174, 176, 177, 182, 187, 189, 193, 194, 200, 205, 208, 210, 219, 224, 225, 227, 236, 237, 245, 251, 253, 267, 271, 272, 280, 283, 290, 292, 294, 296, 304, 306, 308, 310, 314, 319, 321, 323, 327, 331, 335, 336, 339, 353, 354, 356, 358, 361, 365, 367, 368, 369, 370, 378, 382.

[0061] The numbers above correspond to amino acid positions in the NotI protein sequence (SEQ ID NO:90).

[0062] The method for introducing mutants into the enzyme was the same as in the previous examples using inverse PCR followed by DpnI digestion. The treated product was then transformed into *E. coli* containing pACYC-EagIM.

3. Selection of NotI-HF

[0063] Selection of NotI-HF was performed as described in the previous examples. The standard cognate and star activity assays used pXba substrate in NEB 4 buffer and 5% glycerol and NEB ExoI duffer (67 mM Glycine-KOH, pH 9.5, 6.7 mM $MgCl_2$, 10 mM 2-mercaptoethanol) and 37% glycerol respectively. #37(K150A), #44(K176A), #45(R177A), #63(R253A) all showed reduced star activity. K150A was the preferred mutation to reduce star activity. NotI(K150A) was selected as the NotI-HF.

4. Comparison of NotI-HF and WT NotI

[0064] The FIs of NotI-HF and WT NotI were determined separately using pXba substrate in NEB1-4 buffers. The comparison is shown in Figure 17, and the results are listed in Table 16 (below).

Table 16: Comparison of NotI-HF and WT NotI

Buffer	NotI-HF		WT NotI		
	Activity	Fidelity Index	Activity	Fidelity Index	Improvement Factor
NEB1	6%	≥8000	6%	≥8000	ND
NEB2	100%	≥128000	50%	250	≥512
NEB3	1.6%	≥4000	100%	4000	≥1
NEB4	50%	≥64000	12.5%	32	≥2000
ND: Not determinable, for that both FI is an uncertain number over limit.					

[0065] NotI-HF performed best in NEB2, in which the preferred FI was ≥128000; WT NheI performed best in NEB3, in which the preferred FI was 4000. The overall fidelity index improvement factor was ≥128000/4000= ≥32. Engineering NotI not only further improved the FI of NotI, but also changed the optimal buffer.

SEQUENCE LISTING

<110> New England Biolabs, Inc.
Zhu, Zhenyu
Blanchard, Aine
Xu, Shuang-yong
Guan, Shengxi
Wei, Hua
Zhang, Penghua
Sun, Dapeng
Chang, Siu-Hong

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Ile Ala Thr Asn Gly Gln Glu Val Asp Arg Arg Ala Leu Glu Ala Ala
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Arg Ala His Asn Asp Leu Glu Leu Arg Leu Pro Asp Leu Ile Ala Gly
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Gly Arg Val Ser His Leu Arg Asn Ile Glu Tyr Leu Phe Leu Ala Gly
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Lys Val Asn Val Phe Tyr Ile Pro Glu Lys Ala Ser Trp Glu Tyr Leu
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Arg Glu Asn Asn His Asn Leu Cys Phe Val Gly Gln Glu Gln Asn Glu
225                 230                 235                 240

Leu Thr Ile Lys Leu Ala Lys Met Asn Gly Ile Leu His Gly Ile Asn
                245                 250                 255

Pro Glu Ile Arg Gln Gly Asp Ser Leu Leu Asn Asp Arg Tyr Pro Glu
            260                 265                 270

Leu Lys Ala Glu Ile Val Ile Ser Asn Pro Pro Phe Asn Met Lys Asp
        275                 280                 285

Trp Gly Ala Glu Arg Leu Pro Leu Asn Asp Lys Arg Leu Ile Gly Pro
    290                 295                 300

Val Thr Asn Ser Asn Ala Asn Tyr Met Trp Ile Gln His Phe Leu Tyr
305                 310                 315                 320

His Leu Lys Asp Gly Gly Leu Ala Gly Phe Val Ile Ala Asn Gly Ala
                325                 330                 335

Leu Thr Ser Asn Leu Ala Ala Glu Lys Ile Val Arg Lys His Leu Ile
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340 345 350

Asp Asn Asp Tyr Val Asp Cys Val Val Gln Leu Pro Glu Lys Met Phe
355 360 365

Phe Gly Thr Gly Ile Pro Ser Ala Leu Val Phe Leu Ser Lys Asn Arg
370 375 380

Asn Gly Ser Asn Gly His Ala Lys Arg Glu Lys Glu Val Leu Phe Ile
385 390 395 400

Asp Ala Ser Asp Lys Gly Thr Leu Val Gly Lys Lys Asn Lys Ile Phe
405 410 415

Leu Asp Asp Glu Ile Lys Glu Ile Ala Asp Leu Tyr His Ser Phe Lys
420 425 430

Phe Leu Asn Asp Asn Asp Tyr Asn His Ser Gly Phe Tyr Lys Lys Val
435 440 445

Asn Ile Glu Lys Ile Val Glu Asn Asp Tyr Lys Leu Thr Pro Thr Leu
450 455 460

Tyr Val Gly Val Lys Glu Glu Thr Glu Met Glu Lys Pro Phe Arg Glu
465 470 475 480

Met Ile Ile Glu Tyr Lys Ala Ile Leu Glu Gln Gln Phe Glu Glu Ser
485 490 495

Asn Lys Leu Gln Gln Lys Ile Leu Lys Asn Leu Glu Gly Leu Leu
500 505 510

<210> 10
<211> 1179
<212> DNA
<213> Bacillus stearothermophilus X1

<220>
<221> misc_feature
<223> DNA sequence of S.BstXI

<400> 10
atgaaaagta ctttgaagga atataaattg ggtgatatta ccgaagtcgt taatggtgcc 60

actccttcaa ctaaaaagcc tgagtactat gaaaatggta caattccatg gattactcct 120

aaagatttat caggctatta ctttaaatat atatctcatg gtgaacgtaa tataacagag 180

cttggtctaa gaaatagttc agctaagttg ttaccaaaag gaactgtatt attttcctca 240

agagccccaa taggatacgt agcaatagct gataattggt taactacgaa ccagggattt 300

aaaagtttta tatgtaatga ggagattatt tacaatgaat acctttatta ttttcttatt 360

gctaaaaggg attttattga aacatttgcg aatgggagta cgtttaaaga gctttcatca 420

acttctgcaa agaatatacc aatcaatctt cctagtttag aagagcaaaa gaagattgtg 480

acaattttag gggatttgga tagaaagata gaattaaatt ataaaattat tgaaagctta 540

gaaaaaatag cagaaagaac atataaatat tggtttgtcg atgaattaaa tcaagatgaa 600

cagcacatcc gtaatggatg ggaaactgct aaaattggcg atgtggtgga acttttggga 660

gggggaaccc ctaaaacttc ggaaagtaag tattgggaag atggagatat taattggttt 720

actccttcag atttaacaaa aactagacag ctttttgtac gtgattctca aagaaaaata 780

acaattgatg gacttaataa cagtgcagcg aaattaattc ccccttattc cttgttaatg 840

tcaagtagag ctacaattgg cgagttggca attaatcaag aatctgctac tacaaatcaa 900

gggtttattg tattaatacc aaatgaaaaa atttctattt accaattata cttttgggct 960

aaacttaata agagcaaaat tatttcaatg gcaaatggta gtacttttaa agaaattagt 1020

aagcgggatt ttaaatcttt ggagataata ttaccaaaaa atatagacac ttttaattca 1080

attatgcaag attattttag gaaaattgag gagttaattg atgaaataaa aatcttaaaa 1140

accgcaagag ataatttaat tccaaaactt ataaaatga 1179

<210> 11
<211> 392
<212> PRT
<213> Bacillus stearothermophilus X1

<220>
<221> misc_feature
<223> protein sequence for S.BstXI

<400> 11

Met Lys Ser Thr Leu Lys Glu Tyr Lys Leu Gly Asp Ile Thr Glu Val
1 5 10 15

Val Asn Gly Ala Thr Pro Ser Thr Lys Lys Pro Glu Tyr Tyr Glu Asn
20 25 30

Gly Thr Ile Pro Trp Ile Thr Pro Lys Asp Leu Ser Gly Tyr Tyr Phe
35 40 45

Lys Tyr Ile Ser His Gly Glu Arg Asn Ile Thr Glu Leu Gly Leu Arg
50 55 60

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Asn Ser Ser Ala Lys Leu Leu Pro Lys Gly Thr Val Leu Phe Ser Ser
65                  70                  75                  80

Arg Ala Pro Ile Gly Tyr Val Ala Ile Ala Asp Asn Trp Leu Thr Thr
                85                  90                  95

Asn Gln Gly Phe Lys Ser Phe Ile Cys Asn Glu Glu Ile Ile Tyr Asn
            100                 105                 110

Glu Tyr Leu Tyr Tyr Phe Leu Ile Ala Lys Arg Asp Phe Ile Glu Thr
        115                 120                 125

Phe Ala Asn Gly Ser Thr Phe Lys Glu Leu Ser Ser Thr Ser Ala Lys
    130                 135                 140

Asn Ile Pro Ile Asn Leu Pro Ser Leu Glu Glu Gln Lys Lys Ile Val
145                 150                 155                 160

Thr Ile Leu Gly Asp Leu Asp Arg Lys Ile Glu Leu Asn Tyr Lys Ile
                165                 170                 175

Ile Glu Ser Leu Glu Lys Ile Ala Glu Arg Thr Tyr Lys Tyr Trp Phe
            180                 185                 190

Val Asp Glu Leu Asn Gln Asp Glu Gln His Ile Arg Asn Gly Trp Glu
        195                 200                 205

Thr Ala Lys Ile Gly Asp Val Val Glu Leu Leu Gly Gly Gly Thr Pro
    210                 215                 220

Lys Thr Ser Glu Ser Lys Tyr Trp Glu Asp Gly Asp Ile Asn Trp Phe
225                 230                 235                 240

Thr Pro Ser Asp Leu Thr Lys Thr Arg Gln Leu Phe Val Arg Asp Ser
                245                 250                 255

Gln Arg Lys Ile Thr Ile Asp Gly Leu Asn Asn Ser Ala Ala Lys Leu
            260                 265                 270

Ile Pro Pro Tyr Ser Leu Leu Met Ser Ser Arg Ala Thr Ile Gly Glu
        275                 280                 285

Leu Ala Ile Asn Gln Glu Ser Ala Thr Thr Asn Gln Gly Phe Ile Val
    290                 295                 300
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Leu Ile Pro Asn Glu Lys Ile Ser Ile Tyr Gln Leu Tyr Phe Trp Ala
305 310 315 320

Lys Leu Asn Lys Ser Lys Ile Ile Ser Met Ala Asn Gly Ser Thr Phe
325 330 335

Lys Glu Ile Ser Lys Arg Asp Phe Lys Ser Leu Glu Ile Ile Leu Pro
340 345 350

Lys Asn Ile Asp Thr Phe Asn Ser Ile Met Gln Asp Tyr Phe Arg Lys
355 360 365

Ile Glu Glu Leu Ile Asp Glu Ile Lys Ile Leu Lys Thr Ala Arg Asp
370 375 380

Asn Leu Ile Pro Lys Leu Ile Lys
385 390

<210> 12
<211> 770
<212> DNA
<213> planococcus citreus SE-F45

<400> 12
atgaaacagt ttgcagatcc ttttgaaaga agattccttg atgcaattga acatcatctt 60

gatggaattt ctgagaaaat aaaaaaagac tttacacaca aaaacttttt aaaagaattg 120

aatggcctta aaggtgataa agtctatcat gacttaggct ttgataccgc tgaatatact 180

ctggtacgtc ttataggaag aatgagcata agcgttggga gaaggctggg ggagatatac 240

gataaagtcc ctcgttatgt tgctgccgcg cgatttggtc ttcaaccaaa tcaaattgca 300

gaagtatttg atggtcttga gttagatata gctttgcgca atagcctttt gtcagatgat 360

gataaaattc acataaaaaa aataactgaa aagatgtcag gcgaaacata ctcgggaatc 420

ggaatcgaaa ttcgttataa ctttaatcca aatgacagtt cccgtttaag aaaagacgtc 480

gatgtagctt ctaaattgtc ggccgcgggg ttatttcctg tttatttaat atttagctct 540

ctcagtccta ggaatgatgc aatagcccgt cttaaaagag ggggatggag ctttaaacag 600

gggcaggaag ccttagactt ccttaccgaa cttttaggag tggatattgg gtctgtttta 660

tctgacccaa taatagccgc agaaactagg gagaaaacat caaaaattat gaagtctata 720

tttgaatcag aggcattcca atctgttata ccgggagagt ggagtaaact 770

<210> 13
<211> 257
<212> PRT
<213> planococcus citreus SE-F45

<400> 13

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Met Lys Gln Phe Ala Asp Pro Phe Glu Arg Arg Phe Leu Asp Ala Ile
1               5                  10                  15

Glu His His Leu Asp Gly Ile Ser Glu Lys Ile Lys Lys Asp Phe Thr
            20                  25                  30

His Lys Asn Phe Leu Lys Glu Leu Asn Gly Leu Lys Gly Asp Lys Val
        35                  40                  45

Tyr His Asp Leu Gly Phe Asp Thr Ala Glu Tyr Thr Leu Val Arg Leu
    50                  55                  60

Ile Gly Arg Met Ser Ile Ser Val Gly Arg Arg Leu Gly Glu Ile Tyr
65                  70                  75                  80

Asp Lys Val Pro Arg Tyr Val Ala Ala Ala Arg Phe Gly Leu Gln Pro
                85                  90                  95

Asn Gln Ile Ala Glu Val Phe Asp Gly Leu Glu Leu Asp Ile Ala Leu
            100                 105                 110

Arg Asn Ser Leu Leu Ser Asp Asp Asp Lys Ile His Ile Lys Lys Ile
        115                 120                 125

Thr Glu Lys Met Ser Gly Glu Thr Tyr Ser Gly Ile Gly Ile Glu Ile
    130                 135                 140

Arg Tyr Asn Phe Asn Pro Asn Asp Ser Ser Arg Leu Arg Lys Asp Val
145                 150                 155                 160

Asp Val Ala Ser Lys Leu Ser Ala Ala Gly Leu Phe Pro Val Tyr Leu
                165                 170                 175

Ile Phe Ser Ser Leu Ser Pro Arg Asn Asp Ala Ile Ala Arg Leu Lys
            180                 185                 190

Arg Gly Gly Trp Ser Phe Lys Gln Gly Gln Glu Ala Leu Asp Phe Leu
        195                 200                 205

Thr Glu Leu Leu Gly Val Asp Ile Gly Ser Val Leu Ser Asp Pro Ile
    210                 215                 220

Ile Ala Ala Glu Thr Arg Glu Lys Thr Ser Lys Ile Met Lys Ser Ile
225                 230                 235                 240
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Phe Glu Ser Glu Ala Phe Gln Ser Val Ile Pro Gly Glu Trp Ser Lys
245 250 255

Leu

<210> 14
<211> 1347
<212> DNA
<213> planococcus citreus SE-F45

<220>
<221> misc_feature
<223> DNA sequence for M.PciI

<400> 14
atgacaaatt tttcgcactc agctctaacg agctacgatc ttctcgggca tgaaattgtc 60

caagattctg aagctgttag ctcgggtcca tatctggtca gctatgaccc gatccctgta 120

cgtcggtcta cattcctagc tggactgtca gagaacgttc actcgtggtt tcgtctcaca 180

ccaagtttcg gaccggatct agttcgaaca atcatcaaac agatgaatct tgcgccgcac 240

tcacacatcc atgacccttt ctcaggagcc gggactaccg cgattgaggc ttcgttagag 300

ggctatgaag caagctgcgt agaagttaat ccgtttctct acttcgtggg gaaaacatcc 360

atagattggt ctatcaatgc tgatgatgct gcagcgcagc tagaaagcat taaaaataaa 420

tattatagca tgtctgcaac cgctactttg gataacatag ccgacctagg aatagatata 480

ccaaaaatac acaatattca tcggtggtgg agaaacgatg ttcttaaaga tatattagtc 540

ctaaaatctt ctatcagatc ttgcacacaa gataagtatt gttccttttt tgagctagcc 600

ctagctgcag ttctcgttcc agatttgaca aatgtaacgc taggaaaact acaactgcac 660

tttgtaaaca aagacgataa agagataaac gtctggccta catatgaatc tcatgcaaaa 720

aaaatgattc acgacttgtc attaattaat aagcaaaatt tcgaattttt gcccaagatt 780

atttatggtg attcaactca aaaatcaaca tttagcgagg tggcagggat agatgctata 840

ataacatccc ctccgtaccc taataggtac agctatattt ggaatactcg ccctcacctg 900

tacattcttg atatgatttc cgaagcaaaa gaggcttcgc aaatagatcg tagaacgatt 960

ggtggaacat gggggacagc aacttccgaa ttaggaaagg gtatattttc tccaatcaat 1020

gctgtagtca aagacgcgct tgaaggggtt cacgaaagaa tcgccggttc cgatcaactc 1080

atggcaaact atgtaactca ttattttaat cggctctttt tacatataga agctataaaa 1140

ccatcactta atccaaaagc aaagcttgct tatgttgttg ggaactcttg gattaagggc 1200

gaatatgtag ccactgacgt aatcttagca aaaattatcg aaggggcttt gccaggctca 1260

tcaattgatg gtcttcatcg tttccgtcgc cggaacagtg gaaagaatct ctttgaaact 1320

atagtttact ccactctccc ggtataa 1347

<210> 15
<211> 448
<212> PRT
<213> planococcus citreus SE-F45

<220>
<221> misc_feature
<223> Protein sequence for M.PciI

<400> 15

Met Thr Asn Phe Ser His Ser Ala Leu Thr Ser Tyr Asp Leu Leu Gly
1 5 10 15

His Glu Ile Val Gln Asp Ser Glu Ala Val Ser Ser Gly Pro Tyr Leu
20 25 30

Val Ser Tyr Asp Pro Ile Pro Val Arg Arg Ser Thr Phe Leu Ala Gly
35 40 45

Leu Ser Glu Asn Val His Ser Trp Phe Arg Leu Thr Pro Ser Phe Gly
50 55 60

Pro Asp Leu Val Arg Thr Ile Ile Lys Gln Met Asn Leu Ala Pro His
65 70 75 80

Ser His Ile His Asp Pro Phe Ser Gly Ala Gly Thr Thr Ala Ile Glu
85 90 95

Ala Ser Leu Glu Gly Tyr Glu Ala Ser Cys Val Glu Val Asn Pro Phe
100 105 110

Leu Tyr Phe Val Gly Lys Thr Ser Ile Asp Trp Ser Ile Asn Ala Asp
115 120 125

Asp Ala Ala Ala Gln Leu Glu Ser Ile Lys Asn Lys Tyr Tyr Ser Met
130 135 140

Ser Ala Thr Ala Thr Leu Asp Asn Ile Ala Asp Leu Gly Ile Asp Ile
145 150 155 160

Pro Lys Ile His Asn Ile His Arg Trp Trp Arg Asn Asp Val Leu Lys
165 170 175

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Asp Ile Leu Val Leu Lys Ser Ser Ile Arg Ser Cys Thr Gln Asp Lys
            180                 185                 190

Tyr Cys Ser Phe Phe Glu Leu Ala Leu Ala Ala Val Leu Val Pro Asp
        195                 200                 205

Leu Thr Asn Val Thr Leu Gly Lys Leu Gln Leu His Phe Val Asn Lys
    210                 215                 220

Asp Asp Lys Glu Ile Asn Val Trp Pro Thr Tyr Glu Ser His Ala Lys
225                 230                 235                 240

Lys Met Ile His Asp Leu Ser Leu Ile Asn Lys Gln Asn Phe Glu Phe
                245                 250                 255

Leu Pro Lys Ile Ile Tyr Gly Asp Ser Thr Gln Lys Ser Thr Phe Ser
            260                 265                 270

Glu Val Ala Gly Ile Asp Ala Ile Ile Thr Ser Pro Pro Tyr Pro Asn
        275                 280                 285

Arg Tyr Ser Tyr Ile Trp Asn Thr Arg Pro His Leu Tyr Ile Leu Asp
    290                 295                 300

Met Ile Ser Glu Ala Lys Glu Ala Ser Gln Ile Asp Arg Arg Thr Ile
305                 310                 315                 320

Gly Gly Thr Trp Gly Thr Ala Thr Ser Glu Leu Gly Lys Gly Ile Phe
                325                 330                 335

Ser Pro Ile Asn Ala Val Val Lys Asp Ala Leu Glu Gly Val His Glu
            340                 345                 350

Arg Ile Ala Gly Ser Asp Gln Leu Met Ala Asn Tyr Val Thr His Tyr
        355                 360                 365

Phe Asn Arg Leu Phe Leu His Ile Glu Ala Ile Lys Pro Ser Leu Asn
    370                 375                 380

Pro Lys Ala Lys Leu Ala Tyr Val Val Gly Asn Ser Trp Ile Lys Gly
385                 390                 395                 400

Glu Tyr Val Ala Thr Asp Val Ile Leu Ala Lys Ile Ile Glu Gly Ala
                405                 410                 415
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Leu Pro Gly Ser Ser Ile Asp Gly Leu His Arg Phe Arg Arg Arg Asn
420 425 430

Ser Gly Lys Asn Leu Phe Glu Thr Ile Val Tyr Ser Thr Leu Pro Val
435 440 445

<210> 16
<211> 430
<212> PRT
<213> Bacillus sphaericus

<400> 16

Met Arg Arg Leu Ala Lys Asn Ser Arg Asn Asp Ser Tyr Leu Ser Asn
1 5 10 15

Arg Asp Tyr Gln Glu Ile Val Arg Glu Asn Thr Thr Thr Ile Ser Phe
20 25 30

Pro Leu Lys Glu Lys His Thr Leu Thr Leu Thr Lys Lys Ile Gly Leu
35 40 45

Asn Gln Thr Ala Gly Phe Gly Gly Trp Phe Phe Pro Asp Ser Pro Cys
50 55 60

Leu Leu Thr Val Thr Val Leu Ser Ser Phe Gly Thr Lys Val Thr Ser
65 70 75 80

Lys Thr Phe Ser Leu Ser Lys Asp Trp Asn Arg Val Gly Leu Ala Trp
85 90 95

Ile Asn Glu His Ser Ser Asp Thr Met Ser Ile Val Leu Glu Phe Ser
100 105 110

Asp Val Glu Ile Val His Thr Trp Gly Leu Thr Cys Asp Val Phe Asn
115 120 125

Val His Glu Leu Ile Ile Asp Ala Ile Glu Asp Gln Asn Lys Leu Ile
130 135 140

Asp Val Leu Asn Gln Glu His Leu Ser Pro Glu Thr Tyr Tyr Leu Asn
145 150 155 160

His Asp Ser Asp Thr Asp Leu Ile Glu Asn Leu Glu Ser Thr Glu Glu
165 170 175

Ile Lys Ile Val Asn Gln Ser Gln Lys Gln Ile Ser Leu Lys Lys Cys

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                180                 185                 190

Cys Tyr Cys Gln Arg Tyr Met Pro Val Asn Ile Leu Val Arg Ser Asn
        195                 200                 205

Ser Ser Phe His Lys His Lys Ser Lys Lys Thr Gly Phe Gln Asn Glu
    210                 215                 220

Cys Arg Ala Cys Lys Lys Trp Arg Ile Asn Asn Ser Phe Asn Pro Val
225                 230                 235                 240

Arg Thr Lys Asp Gln Leu His Glu Ser Ala Val Ile Thr Arg Glu Lys
                245                 250                 255

Lys Ile Leu Leu Lys Glu Pro Glu Ile Leu Gln Lys Ile Lys Asn Arg
            260                 265                 270

Asn Asn Gly Glu Gly Leu Lys Ser Ile Ile Trp Lys Lys Phe Asp Lys
        275                 280                 285

Lys Cys Phe Asn Cys Glu Lys Glu Leu Thr Ile Glu Glu Val Arg Leu
    290                 295                 300

Asp His Thr Arg Pro Leu Ala Tyr Leu Trp Pro Ile Asp Glu His Ala
305                 310                 315                 320

Thr Cys Leu Cys Glu Lys Cys Asn Asn Thr Lys His Asp Met Phe Pro
                325                 330                 335

Ile Asp Phe Tyr Gln Gly Asp Glu Asp Lys Leu Arg Arg Leu Ala Arg
            340                 345                 350

Ile Thr Gly Leu Asp Tyr Glu Ser Leu Val Lys Arg Asp Val Asn Glu
        355                 360                 365

Val Glu Leu Ala Arg Ile Ile Asn Asn Ile Glu Asp Phe Ala Thr Asn
    370                 375                 380

Val Glu Ala Arg Thr Phe Arg Ser Ile Arg Asn Lys Val Lys Glu Val
385                 390                 395                 400

Arg Pro Asp Thr Asp Leu Phe Glu Ile Leu Lys Ser Lys Asn Ile Asn
                405                 410                 415

Leu Tyr Asn Glu Leu Gln Tyr Glu Leu Leu Thr Arg Lys Asp
            420                 425                 430
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<210> 17
<211> 1245
<212> DNA
<213> Enterobacter aerogenes

<400> 17
gtgaatcaga aaaatgaaaa atcatttatg cgtttgcaat caacctttag cggtggcaaa 60

ggtagtccaa tgcatgattg gtacccatat ttagagggtt attctcccga atttgtgaaa 120

tgcttgattt cacgatttgc tcctaaagcc aaaacaattt tagatccatt ttgtggctct 180

ggaacaacag ccattgtttc cgttttagag ggtttaaata attactattg cgaagtaaac 240

cctttatgcc aatatattat tgaaactaaa ctaatagctt taacattaag cgaagaagaa 300

aaaacaaaat tagtaaatga actttattct atttctaatg aaataactaa tgtactcaaa 360

ccttctgcaa ccgagacaga tctagagaaa tcatttaaat ccgtttttgg taatacgaaa 420

ttttttgagg atcacatatt taaagatata cttagttatc aatgttacat tagctctatc 480

gaagatgaaa atcttaagag acttctgaca atagcaggga ttagatcgtt aatcccttcc 540

tcgttattgg taagacgagg tgatttacga ttcaagacac aaaaagaatt agagaaaggc 600

aaccagggct ttcgctttca tgtacaaaaa agcttagaat taattgccag tgatttatta 660

gacattacgg aaggtagtgg tttagctacc ttcttatgtg atgatgccaa agaaatatct 720

gggaataacc tgattgatgc tgtaataaca agcccgccat atttaaatgg cacaaattat 780

tttagaaata ctaaaattga actttggttt atagggaaat taaagaccaa atcagatcta 840

agacattata gggatttagc tattaccagt ggtattaacg atgtaactaa aggtaaaagc 900

ttatcttcaa ataatactat tatctcagaa ataccattat tatctgaatg tattaaagaa 960

ctaagcataa aagagtatga tagtcgtatt tcaatgatgg ttgaaaacta cttttgggac 1020

atgttcaaat tcttatcaaa actcccaaaa ttactaacta atgatgcgac tatctgtata 1080

gatttaggtg attctgttta ttgtaacgtc tacatcccta cacaagatat tttgaaagaa 1140

atgatgtcaa agttaggttt tgaagagaac gaaagggtca ttcttcgtga acgaaaatcc 1200

cgcaatggaa caaagttagt ccagactgtt caggttttta aatga 1245

<210> 18
<211> 1140
<212> DNA
<213> Enterobacter aerogenes

<400> 18
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tcaggtgaaa tggtaaagag aaattggggc cataactggc actctatgtg ttcataccaa 120

gggaaactta aaccatcaat agctagatct ttaattgata cattcatgcc atcaagtaag 180

ggacgtatat tagatgtctt ctcaggtgtt ggcaccattc ctttcgaagc aagattactt 240

ggtcatactg catatggatt tgatattagt ccagcagcag ttaatatttc acgcgcaaaa 300

ctagaagtta taagtaaaaa tgaaatccaa gaggtaatta ataaattatc tgattttatt 360

gagcaaaaca aaaattcaat agattataac gaacataatt taataaggtt taatggttca 420

attgaatcct attttcatcc tgaaactttt aaggaaatac tgtgtgctcg taaattcttt 480

ttaataaaag gtgaattaaa tgcatctgaa tcgttagtac agtcatgttt attacatatt 540

ttacatggta atcgtccgta tgcattgagt agaaagtccc atcctattac acctttcgcg 600

cctactggag attttatata cagtaattta gttataaagt taatcaaaaa agttgaaaga 660

gtcttgcaaa attctgatgg tatcccagat actggcagca aagtatttta tcaggactct 720

acaaaaagtt ggcctgaaga agtaaataat ttagatgcaa ttataacatc acctccattt 780

tatgatagta cccgtttcta ttcagcaaat tggatgcgat tatggttttc tggttgggaa 840

aaagatgact tccaaacgaa gccaaaagat tttgtggacg aaactcagaa aaaaagcttt 900

gaaatatatg ataatatatt caaacaatct caacaatgct taaaaaaaga tggcgttttt 960

ttaatgcacg ttggcaaaag taaaaaaagt gatatggcag gacaaattgc taaaattggt 1020

agtaattatc ttagccttat agatatattt gacgaaagtg ttgaacattg cgaaagtcac 1080

ggaattaaag acaaaggcac gacaacccat catcagtacc ttgtctttac gaaagattag 1140

<210> 19
<211> 213
<212> PRT
<213> Bacillus amyloliquefaciens H

<400> 19

Met Glu Val Glu Lys Glu Phe Ile Thr Asp Glu Ala Lys Glu Leu Leu
1 5 10 15

Ser Lys Asp Lys Leu Ile Gln Gln Ala Tyr Asn Glu Val Lys Thr Ser
20 25 30

Ile Cys Ser Pro Ile Trp Pro Ala Thr Ser Lys Thr Phe Thr Ile Asn
35 40 45

Asn Thr Glu Lys Asn Cys Asn Gly Val Val Pro Ile Lys Glu Leu Cys
50 55 60

Tyr Thr Leu Leu Glu Asp Thr Tyr Asn Trp Tyr Arg Glu Lys Pro Leu
65 70 75 80

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Asp Ile Leu Lys Leu Glu Lys Lys Lys Gly Gly Pro Ile Asp Val Tyr
                85                  90                  95

Lys Glu Phe Ile Glu Asn Ser Glu Leu Lys Arg Val Gly Met Glu Phe
            100                 105                 110

Glu Thr Gly Asn Ile Ser Ser Ala His Arg Ser Met Asn Lys Leu Leu
        115                 120                 125

Leu Gly Leu Lys His Gly Glu Ile Asp Leu Ala Ile Ile Leu Met Pro
    130                 135                 140

Ile Lys Gln Leu Ala Tyr Tyr Leu Thr Asp Arg Val Thr Asn Phe Glu
145                 150                 155                 160

Glu Leu Glu Pro Tyr Phe Glu Leu Thr Glu Gly Gln Pro Phe Ile Phe
                165                 170                 175

Ile Gly Phe Asn Ala Glu Ala Tyr Asn Ser Asn Val Pro Leu Ile Pro
            180                 185                 190

Lys Gly Ser Asp Gly Met Ser Lys Arg Ser Ile Lys Lys Trp Lys Asp
        195                 200                 205

Lys Val Glu Asn Lys
    210
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<210> 20
<211> 194
<212> PRT
<213> Oceanospirillum kriegii

<400> 20

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Met Lys Ile Lys Arg Ile Glu Val Leu Ile Asn Asn Gly Ser Val Pro
1               5                   10                  15

Gly Ile Pro Met Ile Leu Asn Glu Ile Gln Asp Ala Ile Lys Thr Val
            20                  25                  30

Ser Trp Pro Glu Gly Asn Asn Ser Phe Val Ile Asn Pro Val Arg Lys
        35                  40                  45

Gly Asn Gly Val Lys Pro Ile Lys Asn Ser Cys Met Arg His Leu His
    50                  55                  60

Gln Lys Gly Trp Ala Leu Glu His Pro Val Arg Ile Lys Ala Glu Met
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65 70 75 80

Arg Pro Gly Pro Leu Asp Ala Val Lys Met Ile Gly Gly Lys Ala Phe
85 90 95

Ala Leu Glu Trp Glu Thr Gly Asn Ile Ser Ser Ser His Arg Ala Ile
100 105 110

Asn Lys Met Val Met Gly Met Leu Glu Arg Val Ile Ile Gly Gly Val
115 120 125

Leu Ile Leu Pro Ser Arg Asp Met Tyr Asn Tyr Leu Thr Asp Arg Val
130 135 140

Gly Asn Phe Arg Glu Leu Glu Pro Tyr Phe Ser Val Trp Arg Gln Phe
145 150 155 160

Asn Leu Lys Asp Ala Tyr Leu Ala Ile Val Glu Ile Glu His Asp Ser
165 170 175

Val Asp Ala Gln Val Ser Leu Ile Pro Lys Gly Thr Asp Gly Arg Ala
180 185 190

Ile Arg

<210> 21
<211> 180
<212> PRT
<213> Bacillus amyloliquefaciens H

<220>
<221> MISC_FEATURE
<222> (1)..(180)
<223> Residues 1-180 correspond to residues 22-201 of the protein sequence of BamHI (seq id no. 19)

<400> 21

Ile Gln Gln Ala Tyr Asn Glu Val Lys Thr Ser Ile Cys Ser Pro Ile
1 5 10 15

Trp Pro Ala Thr Ser Lys Thr Phe Thr Ile Asn Asn Thr Glu Lys Asn
20 25 30

Cys Asn Gly Val Val Pro Ile Lys Glu Leu Cys Tyr Thr Leu Leu Glu
35 40 45

```
Asp Thr Tyr Asn Trp Tyr Arg Glu Lys Pro Leu Asp Ile Leu Lys Leu
    50                  55                  60

Glu Lys Lys Lys Gly Gly Pro Ile Asp Val Tyr Lys Glu Phe Ile Glu
65                  70                  75                  80

Asn Ser Glu Leu Lys Arg Val Gly Met Glu Phe Glu Thr Gly Asn Ile
                85                  90                  95

Ser Ser Ala His Arg Ser Met Asn Lys Leu Leu Leu Gly Leu Lys His
            100                 105                 110

Gly Glu Ile Asp Leu Ala Ile Ile Leu Met Pro Ile Lys Gln Leu Ala
        115                 120                 125

Tyr Tyr Leu Thr Asp Arg Val Thr Asn Phe Glu Glu Leu Glu Pro Tyr
    130                 135                 140

Phe Glu Leu Thr Glu Gly Gln Pro Phe Ile Phe Ile Gly Phe Asn Ala
145                 150                 155                 160

Glu Ala Tyr Asn Ser Asn Val Pro Leu Ile Pro Lys Gly Ser Asp Gly
                165                 170                 175

Met Ser Lys Arg
            180
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<210> 22
<211> 177
<212> PRT
<213> Oceanospirillum kriegii

<220>
<221> MISC_FEATURE
<222> (1)..(177)
<223> Residues 1-177 correspond to residues 18-194 of the protein sequence of OkrAI (seq id no. 20)

<400> 22

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Ile Pro Met Ile Leu Asn Glu Ile Gln Asp Ala Ile Lys Thr Val Ser
1               5                   10                  15

Trp Pro Glu Gly Asn Asn Ser Phe Val Ile Asn Pro Val Arg Lys Gly
            20                  25                  30

Asn Gly Val Lys Pro Ile Lys Asn Ser Cys Met Arg His Leu His Gln
        35                  40                  45
```

Lys Gly Trp Ala Leu Glu His Pro Val Arg Ile Lys Ala Glu Met Arg
50 55 60

Pro Gly Pro Leu Asp Ala Val Lys Met Ile Gly Gly Lys Ala Phe Ala
65 70 75 80

Leu Glu Trp Glu Thr Gly Asn Ile Ser Ser Ser His Arg Ala Ile Asn
85 90 95

Lys Met Val Met Gly Met Leu Glu Arg Val Ile Ile Gly Gly Val Leu
100 105 110

Ile Leu Pro Ser Arg Asp Met Tyr Asn Tyr Leu Thr Asp Arg Val Gly
115 120 125

Asn Phe Arg Glu Leu Glu Pro Tyr Phe Ser Val Trp Arg Gln Phe Asn
130 135 140

Leu Lys Asp Ala Tyr Leu Ala Ile Val Glu Ile Glu His Asp Ser Val
145 150 155 160

Asp Ala Gln Val Ser Leu Ile Pro Lys Gly Thr Asp Gly Arg Ala Ile
165 170 175

Arg

<210> 23
<211> 38
<212> DNA
<213> artificial

<220>
<223> primer

<400> 23
attcaacaag catacaatgc agttaaaaca tctattgt 38

<210> 24
<211> 39
<212> DNA
<213> artificial

<220>
<223> primer

<400> 24
acaaatagat gttttaactg cattgtatgc ttgttgaat 39

<210> 25
<211> 39
<212> DNA
<213> artificial

<220>
<223> primer

<400> 25
caagcataca atgaagttgc aacatctatt tgttcacct 39

<210> 26
<211> 39
<212> DNA
<213> artificial

<220>
<223> primer

<400> 26
aggtgaacaa atagatgttg caacttcatt gtatgcttg 39

<210> 27
<211> 39
<212> DNA
<213> artificial

<220>
<223> primer

<400> 27
acgattaaca acaccgaagc aaattgtaac ggtgtagta 39

<210> 28
<211> 40
<212> DNA
<213> artificial

<220>
<223> primer

<400> 28
acgattaaca acaccgaagc aaattgtaac ggtgtagtat 40

<210> 29
<211> 39
<212> DNA
<213> artificial

<220>
<223> primer

<400> 29
aacggtgtag taccaattgc agaactatgt tacacctta 39

<210> 30

<211> 39
<212> DNA
<213> artificial

<220>
<223> primer

<400> 30
taaggtgtaa catagttctg caattggtac tacaccgtt 39

<210> 31
<211> 39
<212> DNA
<213> artificial

<220>
<223> primer

<400> 31
aacccccttg atatacttgc acttgaaaag aaaaaaggt 39

<210> 32
<211> 39
<212> DNA
<213> artificial

<220>
<223> primer

<400> 32
acctttttc ttttcaagtg caagtatatc aaggggttt 39

<210> 33
<211> 36
<212> DNA
<213> artificial

<220>
<223> primer

<400> 33
gatatactta aacttgcaaa gaaaaaaggt ggtccg 36

<210> 34
<211> 39
<212> DNA
<213> artificial

<220>
<223> primer

<400> 34
cggaccacct tttttctttg caagtttaag tatatcaag 39

<210> 35
<211> 39

<212> DNA
<213> artificial

<220>
<223> primer

<400> 35
atacttaaac ttgaaaaggc aaaaggtggt ccgattgat 39

<210> 36
<211> 40
<212> DNA
<213> artificial

<220>
<223> primer

<400> 36
atcaatcgga ccacctttg ccttttcaa gtttaagtat 40

<210> 37
<211> 39
<212> DNA
<213> artificial

<220>
<223> primer

<400> 37
ggtccgattg atgtttatgc agagttcata gaaaacagt 39

<210> 38
<211> 39
<212> DNA
<213> artificial

<220>
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<400> 38
actgttttct atgaactctg cataaacatc aatcggacc 39

<210> 39
<211> 37
<212> DNA
<213> artificial

<220>
<223> primer

<400> 39
atagaaaaac agtgaacttg cacgtgtagg tatggaa 37

<210> 40
<211> 39
<212> DNA

<213> artificial

<220>
<223> primer

<400> 40
aaattccata cctacacgtg caagttcact gttttctat 39

<210> 41
<211> 39
<212> DNA
<213> artificial

<220>
<223> primer

<400> 41
ggaaatatta gttctgccgc acgttcaatg aacaaactt 39

<210> 42
<211> 39
<212> DNA
<213> artificial

<220>
<223> primer

<400> 42
aagtttgttc attgaaacgt gcggcagaac taatattcc 39

<210> 43
<211> 39
<212> DNA
<213> artificial

<220>
<223> primer

<400> 43
aatattagtt ctgcccacgc atcaatgaac aaacttcta 39

<210> 44
<211> 39
<212> DNA
<213> artificial

<220>
<223> primer

<400> 44
tagaagtttg ttcattgatg cgtgggcaga actaatatt 39

<210> 45
<211> 42
<212> DNA
<213> artificial

<220>
<223> primer

<400> 45
gcccaccgtt caatgaacgc acttctatta ggattaaaac at 42

<210> 46
<211> 42
<212> DNA
<213> artificial

<220>
<223> primer

<400> 46
atgttttaat cctaatagaa gtgcggtcat tgaacggtgg gc 42

<210> 47
<211> 39
<212> DNA
<213> artificial

<220>
<223> primer

<400> 47
attatcctta tgcctattgc acaattggcc tattatctt 39

<210> 48
<211> 39
<212> DNA
<213> artificial

<220>
<223> primer

<400> 48
aagataatag gccaattgtg caataggcat aaggataat 39

<210> 49
<211> 39
<212> DNA
<213> artificial

<220>
<223> primer

<400> 49
ttggcctatt atcttacagc acgtgttacc aatttcgag 39

<210> 50
<211> 39
<212> DNA
<213> artificial

<220>
<223> primer

<400> 50
ctcgaaattg gtaacacgtg ctgtaagata ataggccaa 39

<210> 51
<211> 39
<212> DNA
<213> artificial

<220>
<223> primer

<400> 51
gcctattatc ttacagatgc agttaccaat ttcgaggaa 39

<210> 52
<211> 39
<212> DNA
<213> artificial

<220>
<223> primer

<400> 52
ttcctcgaaa ttggtaactg catctgtaag ataataggc 39

<210> 53
<211> 39
<212> DNA
<213> artificial

<220>
<223> primer

<400> 53
cgtgttacca atttcgaggc attagaacct tattttgaa 39

<210> 54
<211> 39
<212> DNA
<213> artificial

<220>
<223> primer

<400> 54
ttcaaaataa ggttctaatg cctcgaaatt ggtaacacg 39

<210> 55
<211> 39
<212> DNA
<213> artificial

<220>

<223> primer

<400> 55
accaatttcg aggaattagc accttatttt gaacttact 39

<210> 56
<211> 39
<212> DNA
<213> artificial

<220>
<223> primer

<400> 56
agtaagttca aaataaggtg ctaattcctc gaaattggt 39

<210> 57
<211> 42
<212> DNA
<213> artificial

<220>
<223> primer

<400> 57
ccttattttg aacttactgc aggacaacca tttatttta tt 42

<210> 58
<211> 42
<212> DNA
<213> artificial

<220>
<223> primer

<400> 58
aataaaaata aatggttgtg ctgcagtaag ttcaaaataa gg 42

<210> 59
<211> 45
<212> DNA
<213> artificial

<220>
<223> primer

<400> 59
tttatttta ttggatttaa tgctgcagct tataattcta atgtc 45

<210> 60
<211> 45
<212> DNA
<213> artificial

<220>
<223> primer

<400> 60
gacattagaa ttataagctg cagcattaaa tccaataaaa ataaa 45

<210> 61
<211> 39
<212> DNA
<213> artificial

<220>
<223> primer

<400> 61
aatgtccctt taattcccgc aggttctgac ggtatgtca 39

<210> 62
<211> 39
<212> DNA
<213> artificial

<220>
<223> primer

<400> 62
tgacataccg tcagaacctg cgggaattaa agggacatt 39

<210> 63
<211> 39
<212> DNA
<213> artificial

<220>
<223> primer

<400> 63
ttaattccca aaggttctgc aggtatgtca aaacgctca 39

<210> 64
<211> 39
<212> DNA
<213> artificial

<220>
<223> primer

<400> 64
tgagcgtttt gacatacctg cagaaccttt gggaattaa 39

<210> 65
<211> 39
<212> DNA
<213> artificial

<220>
<223> primer

<400> 65
tctgacggta tgtcaaaagc atcaattaag aaatggaaa 39

<210> 66
<211> 39
<212> DNA
<213> artificial

<220>
<223> primer

<400> 66
tttccatttc ttaattgatg cttttgacat accgtcaga 39

<210> 67
<211> 54
<212> DNA
<213> artificial

<220>
<223> primer

<400> 67
ggtggtgcat gcggaggtaa ataaatggaa gtagaaaaag agtttattac tgat 54

<210> 68
<211> 51
<212> DNA
<213> artificial

<220>
<223> primer

<400> 68
ggtggtggta ccctatttgt tttcaacttt atctttccat ttcttaattg a 51

<210> 69
<211> 39
<212> DNA
<213> artificial

<220>
<223> primer

<220>
<221> misc_feature
<222> (19)..(21)
<223> n=a, c, g or t

<400> 69
caagcataca atgaagttnn nacatctatt tgttcacct 39

<210> 70
<211> 39
<212> DNA

<213> artificial

<220>
<223> primer

<220>
<221> misc_feature
<222> (19)..(21)
<223> n=a. c. g or t

<400> 70
aggtgaacaa atagatgtnn naacttcatt gtatgcttg 39

<210> 71
<211> 37
<212> DNA
<213> artificial

<220>
<223> primer

<220>
<221> misc_feature
<222> (16)..(18)

<220>
<221> misc_feature
<222> (16)..(18)
<223> n=a, c, g, or t

<400> 71
gatatactta aacttnnnaa gaaaaaaagg tggtccg 37

<210> 72
<211> 39
<212> DNA
<213> artificial

<220>
<223> primer

<220>
<221> misc_feature
<222> (19)..(21)

<220>
<221> misc_feature
<222> (19)..(21)
<223> n=a, c, g, or t

<400> 72
cggaccacct tttttcttnn naagtttaag tatatcaag 39

<210> 73
<211> 54

<212> DNA
<213> artificial

<220>
<223> primer

<400> 73
ggtggtgcat gcggaggtaa ataaatgtct aataaaaaac agtcaaatag gcta 54

<210> 74
<211> 39
<212> DNA
<213> artificial

<220>
<223> primer

<400> 74
ggtggtggta cctcacttag atctaagctg ttcaaacaa 39

<210> 75
<211> 267
<212> PRT
<213> Escherichia coli RY13

<220>
<221> MISC_FEATURE
<222> (1)..(267)
<223> Amino acid residues 1-267 correspond to residues 4-270 of the protein sequence of EcoRI

<400> 75

Lys Lys Gln Ser Asn Arg Leu Thr Glu Gln His Lys Leu Ser Gln Gly
1 5 10 15

Val Ile Gly Ile Phe Gly Asp Tyr Ala Lys Ala His Asp Leu Ala Val
20 25 30

Gly Glu Val Ser Lys Leu Val Lys Lys Ala Leu Ser Asn Glu Tyr Pro
35 40 45

Gln Leu Ser Phe Arg Tyr Arg Asp Ser Ile Lys Lys Thr Glu Ile Asn
50 55 60

Glu Ala Leu Lys Lys Ile Asp Pro Asp Leu Gly Gly Thr Leu Phe Val
65 70 75 80

Ser Asn Ser Ser Ile Lys Pro Asp Gly Gly Ile Val Glu Val Lys Asp
85 90 95

Asp Tyr Gly Glu Trp Arg Val Val Leu Val Ala Glu Ala Lys His Gln

```
                100                 105                 110

Gly Lys Asp Ile Ile Asn Ile Arg Asn Gly Leu Leu Val Gly Lys Arg
        115                 120                 125

Gly Asp Gln Asp Leu Met Ala Ala Gly Asn Ala Ile Glu Arg Ser His
    130                 135                 140

Lys Asn Ile Ser Glu Ile Ala Asn Phe Met Leu Ser Glu Ser His Phe
145                 150                 155                 160

Pro Tyr Val Leu Phe Leu Glu Gly Ser Asn Phe Leu Thr Glu Asn Ile
                165                 170                 175

Ser Ile Thr Arg Pro Asp Gly Arg Val Val Asn Leu Glu Tyr Asn Ser
            180                 185                 190

Gly Ile Leu Asn Arg Leu Asp Arg Leu Thr Ala Ala Asn Tyr Gly Met
        195                 200                 205

Pro Ile Asn Ser Asn Leu Cys Ile Asn Lys Phe Val Asn His Lys Asp
    210                 215                 220

Lys Ser Ile Met Leu Gln Ala Ala Ser Ile Tyr Thr Gln Gly Asp Gly
225                 230                 235                 240

Arg Glu Trp Asp Ser Lys Ile Met Phe Glu Ile Met Phe Asp Ile Ser
                245                 250                 255

Thr Thr Ser Leu Arg Val Leu Gly Arg Asp Leu
            260                 265
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<210> 76
<211> 265
<212> PRT
<213> Rhodopseudomonas sphaeroides

<220>
<221> MISC_FEATURE
<222> (1)..(265)
<223> amino acid residues 1-265 correspond to residues 10-274 of the protein sequence of RsrI

<400> 76

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Lys Gly Gln Ala Leu Arg Leu Gly Ile Gln Gln Glu Leu Gly Gly Gly
1               5                   10                  15
```

```
Pro Leu Ser Ile Phe Gly Ala Ala Ala Gln Lys His Asp Leu Ser Ile
            20                  25                  30

Arg Glu Val Thr Ala Gly Val Leu Thr Lys Leu Ala Glu Asp Phe Pro
        35                  40                  45

Asn Leu Glu Phe Gln Leu Arg Thr Ser Leu Thr Lys Lys Ala Ile Asn
    50                  55                  60

Glu Lys Leu Arg Ser Phe Asp Pro Arg Leu Gly Gln Ala Leu Phe Val
65                  70                  75                  80

Glu Ser Ala Ser Ile Arg Pro Asp Gly Gly Ile Thr Glu Val Lys Asp
                85                  90                  95

Arg His Gly Asn Trp Arg Val Ile Leu Val Gly Glu Ser Lys His Gln
            100                 105                 110

Gly Asn Asp Val Glu Lys Ile Leu Ala Gly Val Leu Gln Gly Lys Ala
        115                 120                 125

Lys Asp Gln Asp Phe Met Ala Ala Gly Asn Ala Ile Glu Arg Met His
    130                 135                 140

Lys Asn Val Leu Glu Leu Arg Asn Tyr Met Leu Asp Glu Lys His Phe
145                 150                 155                 160

Pro Tyr Val Val Phe Leu Gln Gly Ser Asn Phe Ala Thr Glu Ser Phe
                165                 170                 175

Glu Val Thr Arg Pro Asp Gly Arg Val Val Lys Ile Val His Asp Ser
            180                 185                 190

Gly Met Leu Asn Arg Ile Asp Arg Val Thr Ala Ser Ser Leu Ser Arg
        195                 200                 205

Glu Ile Asn Gln Asn Tyr Cys Glu Asn Ile Val Val Arg Ala Gly Ser
    210                 215                 220

Phe Asp His Met Phe Gln Ile Ala Ser Leu Tyr Cys Lys Ala Ala Pro
225                 230                 235                 240

Trp Thr Ala Gly Glu Met Ala Glu Ala Met Leu Ala Val Ala Lys Thr
                245                 250                 255

Ser Leu Arg Ile Ile Ala Asp Asp Leu
```

260 265

<210> 77
<211> 39
<212> DNA
<213> artificial

<220>
<223> primer

<400> 77
gattgggtgg cgcagaaatt tcaaacgggc cagcagtcg 39

<210> 78
<211> 39
<212> DNA
<213> artificial

<220>
<223> primer

<400> 78
cgactgctgg cccgtttgaa atttctgcgc cacccaatc 39

<210> 79
<211> 272
<212> PRT
<213> Agrobacterium gelatinovorum

<400> 79

Met Arg Leu Asp Leu Asp Phe Gly Arg Gly Leu Val Ala His Val Met
1 5 10 15

Leu Asp Asn Val Ser Glu Glu Gln Tyr Gln Gln Ile Ser Asp Tyr Phe
20 25 30

Val Pro Leu Val Asn Lys Pro Lys Leu Lys Ser Arg Asp Ala Ile Gly
35 40 45

Gln Ala Phe Val Met Ala Thr Glu Val Cys Pro Asp Ala Asn Pro Ser
50 55 60

Asp Leu Trp His His Val Leu Tyr Arg Ile Tyr Ile Arg Glu Lys Ile
65 70 75 80

Gly Thr Asp Pro Ser Gln Ser Trp Val Arg Thr Ser Gly Glu Ala Phe
85 90 95

Glu Val Ala Leu Val Glu Arg Tyr Asn Pro Val Leu Ala Arg His Gly
100 105 110

```
Ile Arg Leu Thr Ala Leu Phe Lys Gly Gln Lys Gly Leu Ala Leu Thr
        115                 120                 125

Arg Met Gly Val Ala Asp Arg Val Gly Ser Arg Lys Val Asp Val Met
    130                 135                 140

Ile Glu Lys Gln Gly Gly Gly Arg Ser Pro Asp Ala Glu Gly Phe Gly
145                 150                 155                 160

Val Val Gly Gly Ile His Ala Lys Val Ser Leu Ala Glu Arg Val Ser
                165                 170                 175

Asp Asp Ile Pro Ala Ser Arg Ile Met Met Gly Glu Gly Leu Leu Ser
            180                 185                 190

Val Leu Ser Thr Leu Asp Val Lys Ser Phe Pro Pro Pro His Gly Asp
        195                 200                 205

Leu Val Asn Arg Gly Glu Leu Gly Thr Pro Asp Arg Pro Ser Asp Lys
    210                 215                 220

Arg Asn Tyr Ile Glu Gly His Gly Asp Phe Ser Ala Cys Phe Ser Tyr
225                 230                 235                 240

Asn Leu Arg Thr Pro Pro Ser Asn Ala Thr Thr Pro Ser Gly Arg His
                245                 250                 255

Ile Tyr Val Ser Ala Ser Leu Val Arg Thr Thr Ser Ser Pro Thr Thr
            260                 265                 270
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<210> 80
<211> 358
<212> PRT
<213> Anabaena variabilis

<400> 80

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Met Glu Glu Asp Leu Asp Leu Ser Glu Asn Ile Glu Ala Ala Ser Ala
1               5                   10                  15

Glu Leu Thr Thr Leu Tyr Gln Val Ala Ala Asp Ala Met Lys Asp Tyr
            20                  25                  30

Ile Glu Ile Tyr Leu Ala Leu Ser Lys Gln Ser Asp Gly Phe Ser Asn
        35                  40                  45

Ile Asn Asn Leu Asp Leu Thr Ser Arg Asn Arg Arg Leu Val Val Ile
```

```
    50                  55                  60

His Gly Leu Ser Leu Glu Leu Asp Pro Asp Thr Ser Thr Pro Glu Glu
65                  70                  75                  80

Ile Lys Arg Glu Ala Glu Arg Met Leu Ala Ile Ala Leu Asp Thr Glu
                85                  90                  95

Ser Ala Ile Thr Ala Gly Val Tyr Glu Lys Met Arg Leu Phe Ala Ser
            100                 105                 110

Ser Leu Val Asp Gln Leu Phe Glu Gln Thr Asp Glu Leu Asn Ser Leu
        115                 120                 125

Ser Ser Glu Tyr Leu Ser Ala Asn Pro Gly Phe Leu Pro Phe Phe Gln
    130                 135                 140

Gln Leu Ala Gly Leu Arg Ser Lys Ser Glu Leu Lys Arg Glu Val Gly
145                 150                 155                 160

Asn Ala Ser Asp Asn Ser Ile Ser Lys Ala Val Ala Glu Arg Ile Leu
                165                 170                 175

Glu Arg Ile Ile Arg Asn Leu Arg Ile Arg Thr Phe Ser Lys Glu Lys
            180                 185                 190

Leu Leu Gln Ala Val Glu Pro Thr Leu Glu Gly Ile Val Arg Asp Leu
        195                 200                 205

Val Gly Lys Val Leu Leu Glu Asn Ile Val Ala Asp Ala Leu Ser Asp
    210                 215                 220

Leu Gln Val Pro Phe Met Arg Glu Ser Glu Tyr Gln Ser Leu Lys Gly
225                 230                 235                 240

Val Ile Tyr Asp Phe Arg Ala Asp Phe Val Ile Pro Asp Ala Gln Asn
                245                 250                 255

Pro Ile Ala Phe Ile Glu Val Arg Lys Ser Ser Thr Arg His Ala Ser
            260                 265                 270

Leu Tyr Ala Lys Asp Lys Met Phe Ser Ala Ile Asn Trp Lys Gly Lys
        275                 280                 285

Asn Lys Arg Leu Leu Gly Ile Leu Val Val Glu Gly Pro Trp Thr Arg
    290                 295                 300
```

Glu Thr Leu Arg Val Met Ala Asn Val Phe Asp Tyr Val Thr Pro Leu
305 310 315 320

Thr Arg Val Ser Gln Val Ala Glu Ala Ile Arg Ala Tyr Leu Asp Gly
325 330 335

Asp Lys Thr Arg Leu Lys Trp Leu Val Asn Phe Ser Ile Glu Glu Ala
340 345 350

Asp His Asp Asn Ile Thr
355

<210> 81
<211> 530
<212> PRT
<213> Bacillus stearothermophilus B61

<400> 81

Met Ala Lys Tyr Gly Arg Gly Lys Phe Leu Pro His Gln Asn Tyr Ile
1 5 10 15

Asp Tyr Met His Phe Ile Val Asn His Lys Asn Tyr Ser Gly Met Pro
20 25 30

Asn Ala Ile Gly Glu Asp Gly Arg Ile Asn Trp Gln Val Ser Ser Gly
35 40 45

Lys Thr Thr Ser Phe Tyr Glu Tyr Tyr Gln Ala Arg Phe Glu Trp Trp
50 55 60

Glu Lys Lys Ala Asp Glu Leu Asn Leu Pro Gly Thr Gly Asn Ser Asn
65 70 75 80

Lys Arg Phe Ser Leu Ala Ala Arg Leu Ile His Pro Thr Gly Gln Arg
85 90 95

Pro Cys Arg Leu Cys Gly Lys Tyr Gln Tyr Val Gly Tyr Met Tyr Val
100 105 110

Ser His Asn Leu Tyr Lys Arg Trp Ser Lys Ile Thr Gly Arg Glu Asp
115 120 125

Leu Phe Phe Lys Lys Gln Asn Ile Ile Glu Ala Ala Asn Ile Phe Lys
130 135 140

```
Ser Ile Met Gly Glu Gln Ala Leu Ile Asn Glu Leu Thr Thr Ile Phe
145                 150                 155                 160

Pro Glu Arg Lys Asp Tyr Phe Asn Arg Leu Pro Asn Ile Glu Asp Phe
                165                 170                 175

Phe Val Ser Ser Ser His Ile Lys Asn Asn Gly Asn Tyr Ile Ser Pro
            180                 185                 190

Gly Phe Met Ala Asn Pro Pro Asp Arg Leu Asp Gly Phe His Asp Tyr
        195                 200                 205

Gly Ile Cys Cys Arg Lys Glu Lys Asp Pro Gly Arg His Asp Asp Asn
    210                 215                 220

Met Arg Leu Tyr Asn His Asp Arg Arg Ala Phe Met Trp Trp Ser Glu
225                 230                 235                 240

Gly Asp Trp Ala Leu Ala Asp Ala Leu Tyr Asn Lys Ala Gly Ala Gly
                245                 250                 255

Lys Cys Ala Asp Pro Asp Cys Gln Lys Glu Val Glu Lys Ile Ser Pro
            260                 265                 270

Asp His Val Gly Pro Ile Ser Cys Gly Phe Lys Gln Ile Pro Phe Phe
        275                 280                 285

Lys Pro Leu Cys Ala Ser Cys Asn Ser Ala Lys Asn Arg Arg Phe Ser
    290                 295                 300

Tyr Gln Asp Val Lys Glu Leu Leu Lys Tyr Glu Asn Tyr Thr Gly Asp
305                 310                 315                 320

Ser Val Ala Ser Trp Gln Val Arg Ala Leu Trp Asp Asn Cys Lys His
                325                 330                 335

Leu Val Lys Asn Asp Asp Asp Ser Lys Leu Leu Ser Asn Leu Met Arg
            340                 345                 350

Ser Leu Gln Asp Tyr Tyr Leu Arg Ser Leu Tyr Lys Leu Phe Ser Asn
        355                 360                 365

Gly Phe Ala His Leu Leu Ser Tyr Phe Leu Thr Pro Glu Tyr Ala His
    370                 375                 380

Tyr Lys Ile Thr Phe Glu Gly Leu Asn Thr Ser Thr Leu Glu Tyr Glu
```

385 390 395 400

Arg Tyr Tyr Lys Thr Phe Lys Lys Thr Lys Ser Thr Ser Ser Leu Ala
405 410 415

Ala Arg Ile Val Arg Ile Ala Phe Glu Glu Leu Glu Ile Tyr Asn Ser
420 425 430

Lys Asp Ile Asn Glu Arg Lys Leu Ile Lys Phe Asp Thr Ser Ser Trp
435 440 445

Glu Lys Asp Phe Glu Asn Ile Ile Ser Tyr Ala Thr Lys Asn Leu Ser
450 455 460

Leu Asp Glu Glu Ala Ser Lys Trp Asn Lys Val Leu Thr Asp Lys Asn
465 470 475 480

Leu Ser Ser Thr Glu Lys Asp Lys Lys Ile Ser Ser Leu Leu Glu Asp
485 490 495

Lys Asn Tyr Glu Val Tyr Lys Lys Gln Phe Tyr Ile Leu Lys Asp Leu
500 505 510

Leu Val Glu His Phe Asn Lys Ile Gly Glu Gln Ile Ala Lys Asp Tyr
515 520 525

Met Lys
530

<210> 82
<211> 301
<212> PRT
<213> Enterobacter agglomerans

<400> 82

Met Lys Lys Arg Arg Asp Leu Val Glu Val Phe Gly Tyr Asn Pro Met
1 5 10 15

Asp Leu Ser Pro Glu Val Arg Ala Leu Trp Asn Leu Gly Ala Cys Pro
20 25 30

Phe Leu Asn Lys Glu Cys Ile Lys Ile Asn His Asp Gln Thr Ile Ile
35 40 45

Tyr Gly Thr Cys Ser Val Thr Ser Pro Tyr Gly Asp Val Ile Ile Cys
50 55 60

```
Pro Asn Arg Leu Tyr Ala Asn Asp Tyr Glu Thr Leu His Lys Val Ser
65                  70                  75                  80

Arg Asp Ala Phe Gly Asp Asp Val Pro Phe Leu Thr Tyr Ser Asn Phe
                85                  90                  95

Ile Lys Tyr Arg Ala Thr Tyr Lys Asp Cys Ile Val Ala Leu Gly Lys
            100                 105                 110

Asn Ser Gly Lys Glu Val Gln Val Gly Arg Ala Leu Ser Met Asp Trp
        115                 120                 125

Val Leu Val Arg Ile Thr Asp Gly Glu Leu Lys Glu Tyr Val Gly Val
    130                 135                 140

Glu Ile Gln Ser Ile Asp Ile Thr Gly Asn Tyr Arg Asp Ala Trp His
145                 150                 155                 160

Ala Tyr Lys Asn Leu Lys Pro Ile Asp Ile Ile Asp Asn Leu Pro Thr
                165                 170                 175

Ser Gln His Gly Leu Asn Trp Ala Asn Val His Lys Arg Leu Ile Pro
            180                 185                 190

Gln Ile Ile Arg Lys Gly Val Val Tyr Ser Arg Ser Asn Tyr Val Lys
        195                 200                 205

Lys Gly Leu Tyr Phe Ile Leu Pro Glu Ile Val Tyr Asn Lys Phe Glu
    210                 215                 220

Asp Val Ile Gly Ala Asp Ile Pro Leu Leu Lys Thr Gln Thr Asn Lys
225                 230                 235                 240

Ser Ile Thr Val His Thr Tyr Ser Leu Gly Glu Pro Ala Ala Asn Gly
                245                 250                 255

Glu Gln Arg Lys Leu Ile Ser Glu Arg Glu Ile Ile Phe Asp Leu Asp
            260                 265                 270

Glu Phe Ser Lys Arg Phe Thr Thr Gly Pro Asn Leu Pro Lys Gly Asp
        275                 280                 285

Asp Leu Asp Ala Val Ile Lys Lys Ala Leu Gly Met Met
    290                 295                 300
```

<210> 83
<211> 277
<212> PRT
<213> Escherichia coli RY13

<400> 83

Met Ser Asn Lys Lys Gln Ser Asn Arg Leu Thr Glu Gln His Lys Leu
1 5 10 15

Ser Gln Gly Val Ile Gly Ile Phe Gly Asp Tyr Ala Lys Ala His Asp
20 25 30

Leu Ala Val Gly Glu Val Ser Lys Leu Val Lys Lys Ala Leu Ser Asn
35 40 45

Glu Tyr Pro Gln Leu Ser Phe Arg Tyr Arg Asp Ser Ile Lys Lys Thr
50 55 60

Glu Ile Asn Glu Ala Leu Lys Lys Ile Asp Pro Asp Leu Gly Gly Thr
65 70 75 80

Leu Phe Val Ser Asn Ser Ser Ile Lys Pro Asp Gly Gly Ile Val Glu
85 90 95

Val Lys Asp Asp Tyr Gly Glu Trp Arg Val Val Leu Val Ala Glu Ala
100 105 110

Lys His Gln Gly Lys Asp Ile Ile Asn Ile Arg Asn Gly Leu Leu Val
115 120 125

Gly Lys Arg Gly Asp Gln Asp Leu Met Ala Ala Gly Asn Ala Ile Glu
130 135 140

Arg Ser His Lys Asn Ile Ser Glu Ile Ala Asn Phe Met Leu Ser Glu
145 150 155 160

Ser His Phe Pro Tyr Val Leu Phe Leu Glu Gly Ser Asn Phe Leu Thr
165 170 175

Glu Asn Ile Ser Ile Thr Arg Pro Asp Gly Arg Val Val Asn Leu Glu
180 185 190

Tyr Asn Ser Gly Ile Leu Asn Arg Leu Asp Arg Leu Thr Ala Ala Asn
195 200 205

Tyr Gly Met Pro Ile Asn Ser Asn Leu Cys Ile Asn Lys Phe Val Asn
210 215 220

His Lys Asp Lys Ser Ile Met Leu Gln Ala Ala Ser Ile Tyr Thr Gln
225 230 235 240

Gly Asp Gly Arg Glu Trp Asp Ser Lys Ile Met Phe Glu Ile Met Phe
245 250 255

Asp Ile Ser Thr Thr Ser Leu Arg Val Leu Gly Arg Asp Leu Phe Glu
260 265 270

Gln Leu Thr Ser Lys
275

<210> 84
<211> 245
<212> PRT
<213> Escherichia coli J62 pLG74

<400> 84

Met Ser Leu Arg Ser Asp Leu Ile Asn Ala Leu Tyr Asp Glu Asn Gln
1 5 10 15

Lys Tyr Asp Val Cys Gly Ile Ile Ser Ala Glu Gly Lys Ile Tyr Pro
20 25 30

Leu Gly Ser Asp Thr Lys Val Leu Ser Thr Ile Phe Glu Leu Phe Ser
35 40 45

Arg Pro Ile Ile Asn Lys Ile Ala Glu Lys His Gly Tyr Ile Val Glu
50 55 60

Glu Pro Lys Gln Gln Asn His Tyr Pro Asp Phe Thr Leu Tyr Lys Pro
65 70 75 80

Ser Glu Pro Asn Lys Lys Ile Ala Ile Asp Ile Lys Thr Thr Tyr Thr
85 90 95

Asn Lys Glu Asn Glu Lys Ile Lys Phe Thr Leu Gly Gly Tyr Thr Ser
100 105 110

Phe Ile Arg Asn Asn Thr Lys Asn Ile Val Tyr Pro Phe Asp Gln Tyr
115 120 125

Ile Ala His Trp Ile Ile Gly Tyr Val Tyr Thr Arg Val Ala Thr Arg
130 135 140

```
Lys Ser Ser Leu Lys Thr Tyr Asn Ile Asn Glu Leu Asn Glu Ile Pro
145                 150                 155                 160

Lys Pro Tyr Lys Gly Val Lys Val Phe Leu Gln Asp Lys Trp Val Ile
                165                 170                 175

Ala Gly Asp Leu Ala Gly Ser Gly Asn Thr Thr Asn Ile Gly Ser Ile
            180                 185                 190

His Ala His Tyr Lys Asp Phe Val Glu Gly Lys Gly Ile Phe Asp Ser
        195                 200                 205

Glu Asp Glu Phe Leu Asp Tyr Trp Arg Asn Tyr Glu Arg Thr Ser Gln
    210                 215                 220

Leu Arg Asn Asp Lys Tyr Asn Asn Ile Ser Glu Tyr Arg Asn Trp Ile
225                 230                 235                 240

Tyr Arg Gly Arg Lys
                245
```

<210> 85
<211> 300
<212> PRT
<213> Haemophilus influenzae Rd (exo-mutant)

<400> 85

```
Met Lys Lys Ser Ala Leu Glu Lys Leu Leu Ser Leu Ile Glu Asn Leu
1               5                   10                  15

Thr Asn Gln Glu Phe Lys Gln Ala Thr Asn Ser Leu Ile Ser Phe Ile
            20                  25                  30

Tyr Lys Leu Asn Arg Asn Glu Val Ile Glu Leu Val Arg Ser Ile Gly
        35                  40                  45

Ile Leu Pro Glu Ala Ile Lys Pro Ser Ser Thr Gln Glu Lys Leu Phe
    50                  55                  60

Ser Lys Ala Gly Asp Ile Val Leu Ala Lys Ala Phe Gln Leu Leu Asn
65                  70                  75                  80

Leu Asn Ser Lys Pro Leu Glu Gln Arg Gly Asn Ala Gly Asp Val Ile
                85                  90                  95

Ala Leu Ser Lys Glu Phe Asn Tyr Gly Leu Val Ala Asp Ala Lys Ser
            100                 105                 110
```

```
Phe Arg Leu Ser Arg Thr Ala Lys Asn Gln Lys Asp Phe Lys Val Lys
        115                 120                 125

Ala Leu Ser Glu Trp Arg Glu Asp Lys Asp Tyr Ala Val Leu Thr Ala
    130                 135                 140

Pro Phe Phe Gln Tyr Pro Thr Thr Lys Ser Gln Ile Phe Lys Gln Ser
145                 150                 155                 160

Leu Asp Glu Asn Val Leu Leu Phe Ser Trp Glu His Leu Ala Ile Leu
                165                 170                 175

Leu Gln Leu Asp Leu Glu Glu Thr Asn Ile Phe Pro Phe Glu Gln Leu
            180                 185                 190

Trp Asn Phe Pro Lys Lys Gln Ser Lys Lys Thr Ser Val Ser Asp Ala
        195                 200                 205

Glu Asn Asn Phe Met Arg Asp Phe Asn Lys Tyr Phe Met Asp Leu Phe
    210                 215                 220

Lys Ile Asp Lys Asp Thr Leu Asn Gln Leu Leu Gln Lys Glu Ile Asn
225                 230                 235                 240

Phe Ile Glu Glu Arg Ser Leu Ile Glu Lys Glu Tyr Trp Lys Lys Gln
                245                 250                 255

Ile Asn Ile Ile Lys Asn Phe Thr Arg Glu Glu Ala Ile Glu Ala Leu
            260                 265                 270

Leu Lys Asp Ile Asn Met Ser Ser Lys Ile Glu Thr Ile Asp Ser Phe
        275                 280                 285

Ile Lys Gly Ile Lys Ser Asn Asp Arg Leu Tyr Leu
    290                 295                 300
```

<210> 86
<211> 254
<212> PRT
<213> Haemophilus parainfluenzae

<400> 86

```
Met Lys Tyr Glu Glu Ile Asn Phe Lys Val Pro Val Glu Ser Pro Tyr
1               5                   10                  15
```

```
Tyr Pro Asn Tyr Ser Gln Cys Val Ile Glu Arg Ile Tyr Ser Ile Leu
            20                  25                  30

Arg Asn Gln Lys Asp Met Gly Asp Asp Arg Ile Ile Ile Asn Thr Asn
        35                  40                  45

Leu Lys Lys Gly Leu Pro Leu Glu Asn Ile Asn Lys Ile Ala Gly Pro
    50                  55                  60

Met Ile Glu Ala Trp Ala Glu Glu Val Phe Ser Gly Ile Arg Asp Asn
65                  70                  75                  80

Arg Asp Asn Gln Tyr Asn Leu Ile Asn Val Glu Ala Gln Glu Arg Leu
                85                  90                  95

Gly Ile Ser Asp Ile Ile Leu Gln Phe Gln Val Asn Asn Asn Val Ile
            100                 105                 110

Thr Gly Asn Val Asp Val Lys Ala Thr Ser Asn Asp Ile Pro Asp Ser
        115                 120                 125

Gly Lys Ser Pro Asn Ile Thr Ser Phe Ser Arg Ile Arg Thr Ala Tyr
    130                 135                 140

Val Lys Asp Pro Asn Phe Ile Phe Ile Ile Leu Ser Ile Lys His Ser
145                 150                 155                 160

Val Tyr Val Lys Arg Asn Glu Tyr Thr Asn Leu Met Asp Gly Ile Met
                165                 170                 175

Gln Ile Ile Asp Phe Asn Val Tyr Asp Leu Lys Tyr Ile Ser Asp Ser
            180                 185                 190

Asp Ile Ser Tyr Asn Pro Ala Leu Gly Thr Gly Gln Ile Gln Ile Lys
        195                 200                 205

Asp Ile His Tyr Val Ser Ser Gln Lys Arg Thr Thr Trp Gln Met Cys
    210                 215                 220

Gln Leu Leu Asp Leu Lys Tyr Leu Arg Ser Lys Lys Arg Thr Ile Glu
225                 230                 235                 240

Gln Phe Tyr Asn Glu Ala Lys Arg Asn Lys Trp Ile Lys Asp
                245                 250
```

<210> 87

<211> 218
<212> PRT
<213> Klebsiella pnermoniae OK8

<400> 87

```
Met Asp Val Phe Asp Lys Val Tyr Ser Asp Asp Asn Asn Ser Tyr Asp
1               5                  10                  15

Gln Lys Thr Val Ser Gln Arg Ile Glu Ala Leu Phe Leu Asn Asn Leu
            20                  25                  30

Gly Lys Val Val Thr Arg Gln Gln Ile Ile Arg Ala Ala Thr Asp Pro
        35                  40                  45

Lys Thr Gly Lys Gln Pro Glu Asn Trp His Gln Arg Leu Ser Glu Leu
    50                  55                  60

Arg Thr Asp Lys Gly Tyr Thr Ile Leu Ser Trp Arg Asp Met Lys Val
65                  70                  75                  80

Leu Ala Pro Gln Glu Tyr Ile Met Pro His Ala Thr Arg Arg Pro Lys
                85                  90                  95

Ala Ala Lys Arg Val Leu Pro Thr Lys Glu Thr Trp Glu Gln Val Leu
            100                 105                 110

Asp Arg Ala Asn Tyr Ser Cys Glu Trp Gln Glu Asp Gly Gln His Cys
        115                 120                 125

Gly Leu Val Glu Gly Asp Ile Asp Pro Ile Gly Gly Gly Thr Val Lys
    130                 135                 140

Leu Thr Pro Asp His Met Thr Pro His Ser Ile Asp Pro Ala Thr Asp
145                 150                 155                 160

Val Asn Asp Pro Lys Met Trp Gln Ala Leu Cys Gly Arg His Gln Val
                165                 170                 175

Met Lys Lys Asn Tyr Trp Asp Ser Asn Asn Gly Lys Ile Asn Val Ile
            180                 185                 190

Gly Ile Leu Gln Ser Val Asn Glu Lys Gln Lys Asn Asp Ala Leu Glu
        195                 200                 205

Phe Leu Leu Asn Tyr Tyr Gly Leu Lys Arg
    210                 215
```

<210> 88
<211> 288
<212> PRT
<213> Nocardia corallina

<400> 88

```
Met Ala Thr Ala Pro Gly His Leu Leu Gly Gln Ile Ile Gly Asn Val
1               5                  10                  15

Met Glu Glu Ala Leu Lys Pro Val Leu Gln Glu Met Ala Asp Arg His
            20                  25                  30

Asp Leu Tyr Leu Asp Ser Lys Gly Leu Arg Pro Gly Val Arg Ser Gly
        35                  40                  45

Ala Leu Val Thr Trp Thr Asp Asp Leu Gly Asn Asn His Asp Leu Asp
    50                  55                  60

Phe Val Leu Glu Arg Gly Gly Ser Ala Thr Lys Ala Gly Asn Pro Ala
65                  70                  75                  80

Ala Phe Ile Glu Ala Ala Trp Arg Arg Tyr Thr Lys His Ser Lys Ala
                85                  90                  95

Lys Ala Gln Glu Ile Gln Gly Ala Val Leu Pro Val Leu Ala Ala Trp
            100                 105                 110

Asn Asn Val Lys Pro Thr Pro Ala Ala Val Val Ala Gly Gln Trp Thr
        115                 120                 125

Ala Pro Ser Leu Gln Gln Met Arg Ser Asn Gly Phe Val Val Leu His
    130                 135                 140

Leu His Phe Pro Thr Thr Ala Gln Val Phe Gly Gly Asn Gly Ile Asn
145                 150                 155                 160

Ile Glu Gly Thr Gly Glu Gly Thr Pro Asp Ala Phe Trp Gln Gln Gln
                165                 170                 175

Cys Asp Ala Tyr Thr Ser Lys Ser Glu Ala Asp Lys Asp Ser Leu Ala
            180                 185                 190

Thr Ala Leu Arg Thr Ala His Ala Gln Glu Phe Arg Thr Phe Val Ala
        195                 200                 205

Glu Leu Glu Arg Arg Val Val Arg Ala Ile Asp Tyr Val Val Val Thr
```

210 215 220

Pro Leu His Gly His Gly Ser Gln Tyr Thr Ser Ile Glu Asn Ala Ile
225 230 235 240

Glu Ala Val Arg Thr Tyr Ser Cys Gly Glu Glu Ser Ala Pro Phe Leu
245 250 255

Arg Phe Glu Ile Arg Ile Ser Tyr Thr Asn Gly Asp Val Ile Gln Ala
260 265 270

Thr Phe Gly Ser Ser Ser Asp Ala Ile Glu Phe Leu Asp Thr Phe Asn
275 280 285

<210> 89
<211> 328
<212> PRT
<213> Neisseria mucosa

<400> 89

Met Ser Ser Tyr His Asp Asp Leu Asn Ile Leu Asn Val Asp Phe Asn
1 5 10 15

His Leu Arg Leu Thr Glu Leu Ile Lys Leu Ala Asp Gln Ala Glu Pro
20 25 30

Phe Tyr Leu Trp Val Glu Lys Ile Phe Arg Gln Val Ser Gly Arg Ala
35 40 45

Asp Ser Leu Glu Thr Ile Ile Glu Val Glu Glu Arg Val Val Leu Lys
50 55 60

Met Ala Ile Leu Thr Cys Phe Thr Ser Asp Glu Lys Glu Leu Pro Lys
65 70 75 80

Leu Phe Asn Gly Val Gly Val Pro Tyr Pro His Ile Lys Ala Cys Tyr
85 90 95

Phe Phe Phe Ala Trp Leu Val Arg Asp Ala Ala Thr Gln Arg Leu Asp
100 105 110

Pro Leu Ile Arg Glu Ala Phe Thr Gln Leu Lys Ser Ile His Pro Gln
115 120 125

Met Lys Lys Thr Glu Leu Glu Ser Glu Ile Phe Ser Gln Leu Leu Val
130 135 140

```
Asn Tyr Arg Asn Glu Leu Ile His Phe Ser Trp Pro Val Ile Arg Glu
145                 150                 155                 160

Val Leu Ile Ser Arg Leu Glu Gly Ser Arg Arg Ala Ala Arg Gly Ser
                165                 170                 175

Tyr Leu Glu Leu Phe Val Arg Thr Ala Leu Ala Gln Ser Ile Thr Tyr
            180                 185                 190

Phe Tyr Lys Ile Tyr Gly Asn Tyr Gly Lys Phe Leu Asp Val Lys Ile
        195                 200                 205

His Asp Lys Pro Leu Lys Val Lys Asn Arg Thr Tyr Asp Val Val Ala
    210                 215                 220

Glu Leu Ile Gly Asn Asn His Asn Thr Gln Tyr Leu Ile Leu Pro Val
225                 230                 235                 240

Lys Thr Arg Glu Thr Gln Gly Gly Gly His Ala His Leu Phe Thr Arg
                245                 250                 255

Asp Ile Glu Gln Ser Asn Asn Asp Ile Arg Glu Leu Tyr Pro Asn Ala
            260                 265                 270

Val Ile Ala Pro Val Ile Ile Ala Glu Asn Trp Ser Asp Thr Glu Lys
        275                 280                 285

Asp Leu Glu Asn Val Gly Tyr Asn Asp Ile Phe His Phe Ser Val Asn
    290                 295                 300

Pro Asn Arg Phe Ala Gly Phe Ser Asp Val Glu Gln Ile Arg Leu Asn
305                 310                 315                 320

Arg Leu Val Glu Arg Ile Leu Leu
                325
```

<210> 90
<211> 383
<212> PRT
<213> Nocardia otitidis-caviarum

<400> 90

```
Met Arg Ser Asp Thr Ser Val Glu Pro Glu Gly Ala Asn Phe Ile Ala
1               5                   10                  15

Glu Phe Phe Gly His Arg Val Tyr Pro Glu Val Val Ser Thr Glu Ala
```

```
                20                  25                  30

Ala Arg Asn Asp Gln Ala Thr Gly Thr Cys Pro Phe Leu Thr Ala Ala
        35                  40                  45

Lys Leu Val Glu Thr Ser Cys Val Lys Ala Glu Thr Ser Arg Gly Val
    50                  55                  60

Cys Val Val Asn Thr Ala Val Asp Asn Glu Arg Tyr Asp Trp Leu Val
65                  70                  75                  80

Cys Pro Asn Arg Ala Leu Asp Pro Leu Phe Met Ser Ala Ala Ser Arg
                85                  90                  95

Lys Leu Phe Gly Tyr Gly Pro Thr Glu Pro Leu Gln Phe Ile Ala Ala
            100                 105                 110

Pro Thr Leu Ala Asp Gln Ala Val Arg Asp Gly Ile Arg Glu Trp Leu
        115                 120                 125

Asp Arg Gly Val His Val Val Ala Tyr Phe Gln Glu Lys Leu Gly Gly
    130                 135                 140

Glu Leu Ser Ile Ser Lys Thr Asp Ser Ser Pro Glu Phe Ser Phe Asp
145                 150                 155                 160

Trp Thr Leu Ala Glu Val Glu Ser Ile Tyr Pro Val Pro Lys Ile Lys
                165                 170                 175

Arg Tyr Gly Val Leu Glu Ile Gln Thr Met Asp Phe His Gly Ser Tyr
            180                 185                 190

Lys His Ala Val Gly Ala Ile Asp Ile Ala Leu Val Glu Gly Ile Asp
        195                 200                 205

Phe His Gly Trp Leu Pro Thr Pro Ala Gly Arg Ala Ala Leu Ser Lys
    210                 215                 220

Lys Met Glu Gly Pro Asn Leu Ser Asn Val Phe Lys Arg Thr Phe Tyr
225                 230                 235                 240

Gln Met Ala Tyr Lys Phe Ala Leu Ser Gly His Gln Arg Cys Ala Gly
                245                 250                 255

Thr Gly Phe Ala Ile Pro Gln Ser Val Trp Lys Ser Trp Leu Arg His
            260                 265                 270
```

Leu Ala Asn Pro Thr Leu Ile Asp Asn Gly Asp Gly Thr Phe Ser Leu
275 280 285

Gly Asp Thr Arg Asn Asp Ser Glu Asn Ala Trp Ile Phe Val Phe Glu
290 295 300

Leu Asp Pro Asp Thr Asp Ala Ser Pro Arg Pro Leu Ala Pro His Leu
305 310 315 320

Glu Ile Arg Val Asn Val Asp Thr Leu Ile Asp Leu Ala Leu Arg Glu
325 330 335

Ser Pro Arg Ala Ala Leu Gly Pro Ser Gly Pro Val Ala Thr Phe Thr
340 345 350

Asp Lys Val Glu Ala Arg Met Leu Arg Phe Trp Pro Lys Thr Arg Arg
355 360 365

Arg Arg Ser Thr Thr Pro Gly Gly Gln Arg Gly Leu Phe Asp Ala
370 375 380

<210> 91
<211> 326
<212> PRT
<213> Providencia stuartii 164

<400> 91

Met Lys Glu Leu Lys Leu Lys Glu Ala Lys Glu Ile Leu Lys Ala Leu
1 5 10 15

Gly Leu Pro Pro Gln Gln Tyr Asn Asp Arg Ser Gly Trp Val Leu Leu
20 25 30

Ala Leu Ala Asn Ile Lys Pro Glu Asp Ser Trp Lys Glu Ala Lys Ala
35 40 45

Pro Leu Leu Pro Thr Val Ser Ile Met Glu Phe Ile Arg Thr Glu Tyr
50 55 60

Gly Lys Asp Tyr Lys Pro Asn Ser Arg Glu Thr Ile Arg Arg Gln Thr
65 70 75 80

Leu His Gln Phe Glu Gln Ala Arg Ile Val Asp Arg Asn Arg Asp Leu
85 90 95

```
Pro Ser Arg Ala Thr Asn Ser Lys Asp Asn Asn Tyr Ser Leu Asn Gln
            100                 105                 110

Val Ile Ile Asp Ile Leu His Asn Tyr Pro Asn Gly Asn Trp Lys Glu
        115                 120                 125

Leu Ile Gln Gln Phe Leu Thr His Val Pro Ser Leu Gln Glu Leu Tyr
    130                 135                 140

Glu Arg Ala Leu Ala Arg Asp Arg Ile Pro Ile Lys Leu Leu Asp Gly
145                 150                 155                 160

Thr Gln Ile Ser Leu Ser Pro Gly Glu His Asn Gln Leu His Ala Asp
                165                 170                 175

Ile Val His Glu Phe Cys Pro Arg Phe Val Gly Asp Met Gly Lys Ile
            180                 185                 190

Leu Tyr Ile Gly Asp Thr Ala Ser Ser Arg Asn Glu Gly Gly Lys Leu
        195                 200                 205

Met Val Leu Asp Ser Glu Tyr Leu Lys Lys Leu Gly Val Pro Pro Met
    210                 215                 220

Ser His Asp Lys Leu Pro Asp Val Val Val Tyr Asp Glu Lys Arg Lys
225                 230                 235                 240

Trp Leu Phe Leu Ile Glu Ala Val Thr Ser His Gly Pro Ile Ser Pro
                245                 250                 255

Lys Arg Trp Leu Glu Leu Glu Ala Ala Leu Ser Ser Cys Thr Val Gly
            260                 265                 270

Lys Val Tyr Val Thr Ala Phe Pro Thr Arg Thr Glu Phe Arg Lys Asn
        275                 280                 285

Ala Ala Asn Ile Ala Trp Glu Thr Glu Val Trp Ile Ala Asp Asn Pro
    290                 295                 300

Asp His Met Val His Phe Asn Gly Asp Arg Phe Leu Gly Pro His Asp
305                 310                 315                 320

Lys Lys Pro Glu Leu Ser
                325
```

<210> 92

<211> 157
<212> PRT
<213> Proteus vulgaris

<400> 92

Met Ser His Pro Asp Leu Asn Lys Leu Leu Glu Leu Trp Pro His Ile
1 5 10 15

Gln Glu Tyr Gln Asp Leu Ala Leu Lys His Gly Ile Asn Asp Ile Phe
20 25 30

Gln Asp Asn Gly Gly Lys Leu Leu Gln Val Leu Leu Ile Thr Gly Leu
35 40 45

Thr Val Leu Pro Gly Arg Glu Gly Asn Asp Ala Val Asp Asn Ala Gly
50 55 60

Gln Glu Tyr Glu Leu Lys Ser Ile Asn Ile Asp Leu Thr Lys Gly Phe
65 70 75 80

Ser Thr His His His Met Asn Pro Val Ile Ile Ala Lys Tyr Arg Gln
85 90 95

Val Pro Trp Ile Phe Ala Ile Tyr Arg Gly Ile Ala Ile Glu Ala Ile
100 105 110

Tyr Arg Leu Glu Pro Lys Asp Leu Glu Phe Tyr Tyr Asp Lys Trp Glu
115 120 125

Arg Lys Trp Tyr Ser Asp Gly His Lys Asp Ile Asn Asn Pro Lys Ile
130 135 140

Pro Val Lys Tyr Val Met Glu His Gly Thr Lys Ile Tyr
145 150 155

<210> 93
<211> 358
<212> PRT
<213> Streptomyces achromogenes

<400> 93

Met Gly Ile Thr Ile Lys Lys Ser Thr Ala Glu Gln Val Leu Arg Lys
1 5 10 15

Ala Tyr Glu Ala Ala Ala Ser Asp Asp Val Phe Leu Glu Asp Trp Ile
20 25 30

```
Phe Leu Ala Thr Ser Leu Arg Glu Val Asp Ala Pro Arg Thr Tyr Thr
        35                  40                  45

Ala Ala Leu Val Thr Ala Leu Leu Ala Arg Ala Cys Asp Asp Arg Val
    50                  55                  60

Asp Pro Arg Ser Ile Lys Glu Lys Tyr Asp Asp Arg Ala Phe Ser Leu
65                  70                  75                  80

Arg Thr Leu Cys His Gly Val Val Val Pro Met Ser Val Glu Leu Gly
                85                  90                  95

Phe Asp Leu Gly Ala Thr Gly Arg Glu Pro Ile Asn Asn Gln Pro Phe
            100                 105                 110

Phe Arg Tyr Asp Gln Tyr Ser Glu Ile Val Arg Val Gln Thr Lys Ala
        115                 120                 125

Arg Pro Tyr Leu Asp Arg Val Ser Ser Ala Leu Ala Arg Val Asp Glu
    130                 135                 140

Glu Asp Tyr Ser Thr Glu Glu Ser Phe Arg Ala Leu Val Ala Val Leu
145                 150                 155                 160

Ala Val Cys Ile Ser Val Ala Asn Lys Lys Gln Arg Val Ala Val Gly
                165                 170                 175

Ser Ala Ile Val Glu Ala Ser Leu Ile Ala Glu Thr Gln Ser Phe Val
            180                 185                 190

Val Ser Gly His Asp Val Pro Arg Lys Leu Gln Ala Cys Val Ala Ala
        195                 200                 205

Gly Leu Asp Met Val Tyr Ser Glu Val Val Ser Arg Arg Ile Asn Asp
    210                 215                 220

Pro Ser Arg Asp Phe Pro Gly Asp Val Gln Val Ile Leu Asp Gly Asp
225                 230                 235                 240

Pro Leu Leu Thr Val Glu Val Arg Gly Lys Ser Val Ser Trp Glu Gly
                245                 250                 255

Leu Glu Gln Phe Val Ser Ser Ala Thr Tyr Ala Gly Phe Arg Arg Val
            260                 265                 270

Ala Leu Met Val Asp Ala Ala Ser His Val Ser Leu Met Ser Ala Asp
```

275 280 285

Asp Leu Thr Ser Ala Leu Glu Arg Lys Tyr Glu Cys Ile Val Lys Val
290 295 300

Asn Glu Ser Val Ser Ser Phe Leu Arg Asp Val Phe Val Trp Ser Pro
305 310 315 320

Arg Asp Val His Ser Ile Leu Ser Ala Phe Pro Glu Ala Met Tyr Arg
325 330 335

Arg Met Ile Glu Ile Glu Val Arg Glu Pro Glu Leu Asp Arg Trp Ala
340 345 350

Glu Ile Phe Pro Glu Thr
355

<210> 94
<211> 315
<212> PRT
<213> Streptomyces albus G

<400> 94

Met Ile Asn Ala Asp Lys Pro His Arg Trp Asn Asp Asp Val Gln Ala
1 5 10 15

Ser Val Arg Leu Tyr Asn Gln Trp Phe Leu Asp Ala Ala Pro Lys Ala
20 25 30

Tyr Arg Asp Thr Arg Gln Leu Thr Ile Asp Glu Val Glu Gln Ala Phe
35 40 45

Gln Arg Thr Ala Asn Met Thr Ser Ile Thr Pro Glu Val Leu Lys Ala
50 55 60

His Pro Lys Thr Leu Ala Thr Leu Arg Met Ser Thr Ala Pro Pro Ile
65 70 75 80

Ala Arg Asp Arg Leu Val Gly Leu Ser His Gly Ser Lys Ser Leu Leu
85 90 95

Asp Thr Met Glu Lys Gly Lys Leu Pro Pro Arg Met Lys Gly Asp Val
100 105 110

Leu Asp Thr His Leu Ala Lys Met Cys Ala Val Leu Thr Asp Leu Leu
115 120 125

```
Asp Leu Asp Leu Phe His Trp Tyr Pro Thr Gly Glu Pro Ala Glu Pro
    130                 135                 140

Arg Gln Arg Glu Leu Ala Ala Thr Val Val Ala Asp Arg Leu Cys Gly
145                 150                 155                 160

Ala Ile Ala Asp Pro Ile Val Arg Asn Ala Gln Glu Arg Arg Gln Leu
                165                 170                 175

Ala Leu Ile Glu Glu Trp Leu Leu Ala Arg Gly Tyr Thr Lys Lys Thr
            180                 185                 190

His Ser Ala Ser Leu Pro Leu Asn Thr Met Gln Pro Gly Thr Phe Ser
        195                 200                 205

Phe Arg Gln Asn Val Val Val Gly Ser Asp Leu Pro Val Asn Ile Pro
    210                 215                 220

Val Asp Ala Val Ile Gln Pro His Thr Pro His Ser His Lys Leu Pro
225                 230                 235                 240

Ile Leu Ile Glu Ala Lys Ser Ala Gly Asp Phe Thr Asn Thr Asn Lys
                245                 250                 255

Arg Arg Lys Glu Glu Ala Thr Lys Ile His Gln Leu Gln Leu Lys Tyr
            260                 265                 270

Gly Asn Glu Ile Ser Leu Thr Leu Phe Leu Cys Gly Tyr Phe Asn Thr
        275                 280                 285

Gly Tyr Leu Gly Tyr Ser Ala Ala Glu Gly Leu Asp Trp Val Trp Glu
    290                 295                 300

His Arg Ile Asp Asp Leu Glu Ala Ala Gly Ala
305                 310                 315
```

<210> 95
<211> 432
<212> PRT
<213> Saccharopolyspora sp.

<400> 95

```
Met Arg Arg Leu Ala Thr Gln Arg Arg Glu Asp Ala Tyr Lys Ser Asn
1               5                   10                  15

Arg Asp Tyr Gln Thr Val His Glu Ala Gln Ser Leu Arg Val Asn Ser
```

```
                  20                  25                  30

Thr Asp Asp Asp Asn Leu Ser Leu Phe Leu Leu Lys Asp Ile Ser Pro
        35                  40                  45

Arg Glu Asp Ser Lys Asn Ile Val Gly Phe Gly Gly Phe Val Lys Pro
    50                  55                  60

Glu Ile Ala Thr Thr Met Ala Leu Thr Leu Thr Thr Asp Ile Asp Lys
65                  70                  75                  80

Gln Ile Lys Ser Val Pro Leu Ser Ser Asn Trp Asn Arg Ile Ser Ile
                85                  90                  95

Val Ala Lys Phe Ala Ser Asn Pro Ser Val Ser Ile Thr Leu Gly Phe
            100                 105                 110

Asp Gln Thr Pro Trp Val Asp Phe Trp Gly Ile Asn Ser Asp Asp Ile
        115                 120                 125

Gly Leu Ser Phe Val Ser Asp Ala Val Pro Leu Glu Met Ser Met Ile
    130                 135                 140

Asp Ser Ile His Ile Ala Pro Glu Thr Leu Tyr Leu Asp His Ser Ser
145                 150                 155                 160

Ala Cys Leu Leu Asp Ile Asp Pro Val Glu Ser Thr Arg Phe Lys Thr
                165                 170                 175

Gly His Gly Asp Pro Leu Ser Leu Lys Lys Cys Ser Tyr Cys Gly Arg
            180                 185                 190

Leu Leu Pro Ile Asp Leu Glu Arg Pro Gly Lys Leu Ser Phe His Lys
        195                 200                 205

His Arg Ala Lys Ile Thr Asn His Gln Asn Glu Cys Arg Ser Cys Lys
    210                 215                 220

Lys Trp Arg Ile Asn Asn Ser Phe Asn Pro Met Arg Thr Ile Asp Gln
225                 230                 235                 240

Leu Asn Glu Ser Ala Leu Ile Thr Arg Glu Arg Lys Ile Phe Leu Gln
                245                 250                 255

Glu Pro Glu Ile Leu Gln Glu Ile Lys Asp Arg Thr Gly Ala Gly Leu
            260                 265                 270
```

Lys Ser Gln Val Trp Glu Arg Phe His Arg Lys Cys Phe Asn Cys Arg
275 280 285

Lys Asp Leu Lys Leu Ser Glu Val Gln Leu Asp His Thr Arg Pro Leu
290 295 300

Ala Tyr Leu Trp Pro Ile Asp Glu His Ala Thr Cys Leu Cys Ala Gln
305 310 315 320

Cys Asn Asn Thr Lys Lys Asp Arg Phe Pro Val Asp Phe Tyr Ser Glu
325 330 335

Gln Gln Ile Arg Glu Leu Ser Asp Ile Cys Gly Leu Pro Tyr Gln Asp
340 345 350

Leu Cys Ala Arg Ser Leu Asn Leu Asp Gln Leu Asp Arg Ile Glu Arg
355 360 365

Asn Ile Ala Glu Phe Ser Lys Glu Trp Asp Val Arg Thr Phe Ala Ser
370 375 380

Thr Ala Arg Arg Ile Ser Glu Val Tyr Pro Ala Arg Asp Leu Phe Glu
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Thr Leu Lys Lys Glu Ser Glu Ser Ala Tyr Asn Lys Ile Ile Glu Lys
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Leu Lys Glu Arg Pro Asp Ala Leu Leu Asp Glu Ala Leu Pro Leu Asp
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<213> Streptomyces species Bf-61

<400> 96

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Val Arg Ser Ala Val Thr Leu Leu Ala Leu Ala Gly Leu Lys Pro Gly
35 40 45

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Asp Arg Trp Val Asp Ser Thr Thr Pro Arg Leu Gly Val Gln Lys Ile
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Met Asp Trp Ser Gly Glu His Trp Ala Lys Pro Tyr Ala Thr Gly Ser
65                  70                  75                  80

Arg Glu Asp Phe Arg Lys Lys Thr Leu Arg Gln Trp Val Asp Asn Gly
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Phe Ala Val Leu Asn Ala Asp Asn Leu Asn Ile Ala Thr Asn Ser Gln
            100                 105                 110

Leu Asn Glu Tyr Cys Leu Ser Asp Glu Ala Leu Gln Ala Leu Arg Ala
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Tyr Gly Thr Glu Gly Phe Glu Glu Ser Leu Val Val Phe Leu Asp Glu
    130                 135                 140

Ala Ser Lys Ala Val Lys Ala Arg Ala Glu Ala Leu Gln Ala Ala Met
145                 150                 155                 160

Ile Ser Val Asp Leu Pro Gly Gly Glu Glu Phe Leu Leu Ser Pro Ala
                165                 170                 175

Gly Gln Asn Pro Leu Leu Lys Lys Met Val Glu Glu Phe Val Pro Arg
            180                 185                 190

Phe Ala Pro Arg Ser Thr Val Leu Tyr Leu Gly Asp Thr Arg Gly Lys
        195                 200                 205

His Ser Leu Phe Glu Arg Glu Ile Phe Glu Glu Val Leu Gly Leu Thr
    210                 215                 220

Phe Asp Pro His Gly Arg Met Pro Asp Leu Ile Leu His Asp Glu Val
225                 230                 235                 240

Arg Gly Trp Leu Phe Leu Met Glu Ala Val Lys Ser Lys Gly Pro Phe
                245                 250                 255

Asp Glu Glu Arg His Arg Ser Leu Gln Glu Leu Phe Val Thr Pro Ser
            260                 265                 270

Ala Gly Leu Ile Phe Val Asn Cys Phe Glu Asn Arg Glu Ser Met Arg
        275                 280                 285

Gln Trp Leu Pro Glu Leu Ala Trp Glu Thr Glu Ala Trp Val Ala Glu
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Asp Pro Asp His Leu Ile His Leu Asn Gly Ser Arg Phe Leu Gly Pro
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Tyr Glu Arg

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<212> PRT
<213> Streptomyces caespitosus

<400> 97

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35 40 45

Leu Ser Glu Gln Pro Leu Ser Gly Asp Thr Leu Val Asp Ala Val Leu
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Ala Ser Trp Glu Ser Ile Phe Glu Ser Arg Leu Gly Ser Gly Phe His
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Ile Gly Thr Gln Ile Arg Pro Thr Pro Gln Ile Met Gly Phe Leu Leu
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His Ala Leu Ile Pro Leu Glu Leu Ala Asn Gly Asp Pro Ser Trp Arg
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Ala Asp Leu Asn Ser Ser Glu Lys Asp Leu Val Tyr Gln Pro Asp His
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Lys Tyr Ser Ile Glu Met Lys Thr Ser Ser His Lys Asp Gln Ile Phe
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Gly Asn Arg Ser Phe Gly Val Glu Asn Pro Gly Lys Gly Lys Lys Ala
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Lys Asp Gly Tyr Tyr Val Ala Val Asn Phe Glu Lys Trp Ser Asp Ala
165 170 175

Pro Gly Arg Leu Pro Arg Ile Arg Thr Ile Arg Tyr Gly Trp Leu Asp
180 185 190

His Thr Asp Trp Val Ala Gln Lys Ser Gln Thr Gly Gln Gln Ser Ser
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Leu Pro Ala Val Val Ser Asn Thr Gln Leu Leu Ala Ile His Thr Gly
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Gly Gln Arg
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<213> Streptomyces phaeochromogenes

<400> 98

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35 40 45

Tyr Ser His Pro Pro Arg Ala Ala Arg Asp Leu Lys Asp Ala Gly Ala
50 55 60

Gly Val Val Thr Ile Met Val Lys Gly Pro Asp Gly Arg Arg Met Ala
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Ser Tyr Ala Phe Asn Gly Lys Ala Asn Glu Asp Gly Ala Gly Arg Val
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Val Ile Pro Lys Ala Phe Gly Glu Ala Leu Lys Arg Ala His Gly Gly
100 105 110

Lys Cys Ala Val Cys Tyr Gly Asp Phe Ser Glu Arg Glu Leu Gln Cys
115 120 125

Asp His Arg Val Pro Phe Ala Ile Ala Gly Asp Lys Pro Lys Leu Val
130 135 140

Gln Glu Asp Phe Met Pro Leu Cys Ala Ser Asp Asn Arg Ala Lys Ser

145 150 155 160

Trp Ser Cys Glu Asn Cys Pro Asn Trp Glu Leu Lys Asp Glu Asp Thr
165 170 175

Cys Arg Ser Cys Phe Trp Ala Ser Pro Glu Asn Tyr Thr His Val Ser
180 185 190

Thr Arg Pro Glu Arg Arg Ile Asn Leu Leu Phe Gln Gly Asp Glu Val
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Glu Ile Phe Asp Ala Leu Lys Asn Ala Ala Ala Asn Glu Gly Val Ser
210 215 220

Leu Thr Glu Ala Thr Lys Arg Lys Leu Ala Asp
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<213> Sphaerotilus species

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Thr Met Arg Leu Ile Gly Asn Lys Thr His Gly Asp Leu Ala Glu Ile
35 40 45

Ala Ile Ser Glu Phe Ile Asn Gln Tyr Met Tyr Asp Phe Lys Ser Ile
50 55 60

His Val Gly Lys Asp Leu Tyr Arg Ala Lys Ser Lys Glu Glu Asp Ile
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Thr Val Glu Asn Glu Ile Thr Lys Glu Lys Phe Pro Ile Ser Leu Lys
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Ala Tyr Gly Asp Gly Pro Leu Gln Leu Ser Thr Asp Lys Asn Phe Leu
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Met Tyr Pro Leu Leu Glu Glu Ile Gly Ala Phe Ile Asn Ala Lys Glu
115 120 125

Lys Ile Glu Glu Ile Phe Ala Asn Glu Ala Phe Ser Cys Phe Ser Glu
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Ile Asn Val Leu Pro Leu Ile Tyr Asp Glu Lys Arg Gln Arg Cys Asn
145 150 155 160

Ile Leu Val Phe Asp Ala Ala Arg Ala Arg Ala Glu Thr Ala Tyr Ile
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Arg Lys Glu Thr Glu Gly Ser Gly Arg Lys His Pro Ala Tyr Arg Phe
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Phe Asp Lys Asn Lys Asn Tyr Ile Cys Glu Val Arg Tyr Gly Asn Ala
195 200 205

Ala Ala Asn Ala Leu Gln Arg Gly Leu Trp Thr Asn Thr Lys Asn Ala
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Thr Ser Phe Phe Asp Ser Val Thr Asn Gly Trp Val Asp Tyr Ser His
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Asn Leu Val Leu Val Lys Leu Leu Ser His Ala Leu Val Ser Ser Arg
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Claims

1. A composition **characterized in that** it comprises: a restriction endonuclease enzyme having at least one artificially introduced mutation and an overall fidelity index (FI) improvement factor of at least 2, the restriction endonuclease being capable of cleaving a substrate with at least a similar cleavage activity to that of the restriction endonuclease absent the artificially introduced mutation, in a predetermined buffer, wherein the artificially introduced mutation is the product of at least one of a targeted mutation, saturation mutagenesis, or a mutation introduced through a PCR amplification procedure, and wherein the restriction endonuclease absent the artificially introduced mutation is NotI and the artificially introduced mutation is selected from: K176A; R177A; R253A; and K150A.

2. A composition according to claim 1 wherein at least one of the artificially introduced mutations is a replacement of a naturally occurring residue with an oppositely charged residue at a target site in the restriction endonuclease.

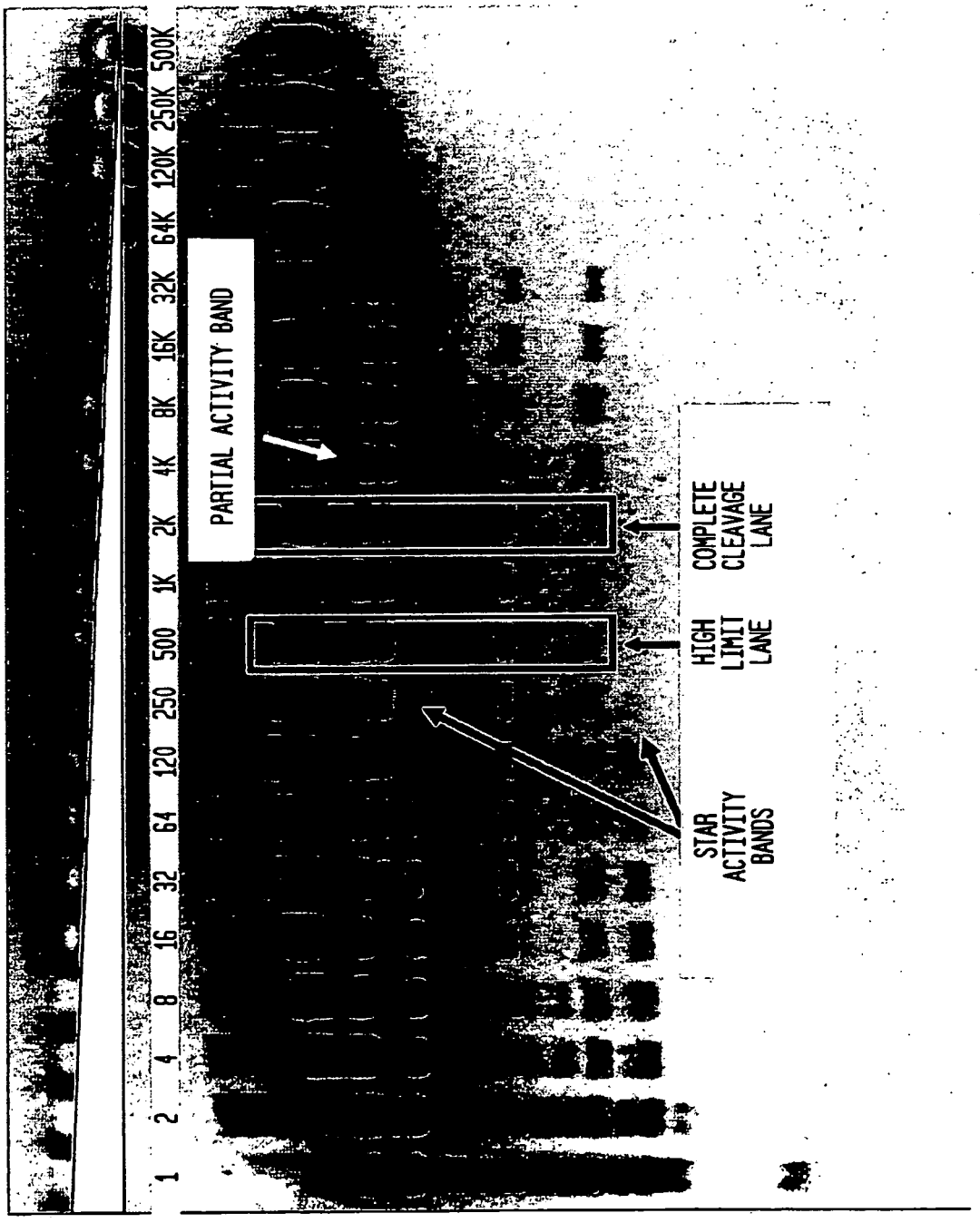
3. A composition according to claim 1 wherein at least one of the artificially introduced mutations is a replacement of a naturally occurring residue with a residue selected from a phenylalanine and an alanine at a target site in the restriction endonuclease.

4. A DNA molecule **characterized in that** it encodes a composition according to claim 1.

5. A vector **characterized in that** it contains a DNA molecule according to claim 4.

6. A host cell **characterized in that** it contains a DNA molecule according to claim 4 or a vector according to claim 5.

FIG. 1


1
2
4
8
16
32
64
120
250
500
1K
2K
4K
8K
16K
32K
64K
120K
250K
500K
PARTIAL ACTIVITY BAND
COMPLETE CLEAVAGE LANE
HIGH LIMIT LANE
STAR ACTIVITY BANDS

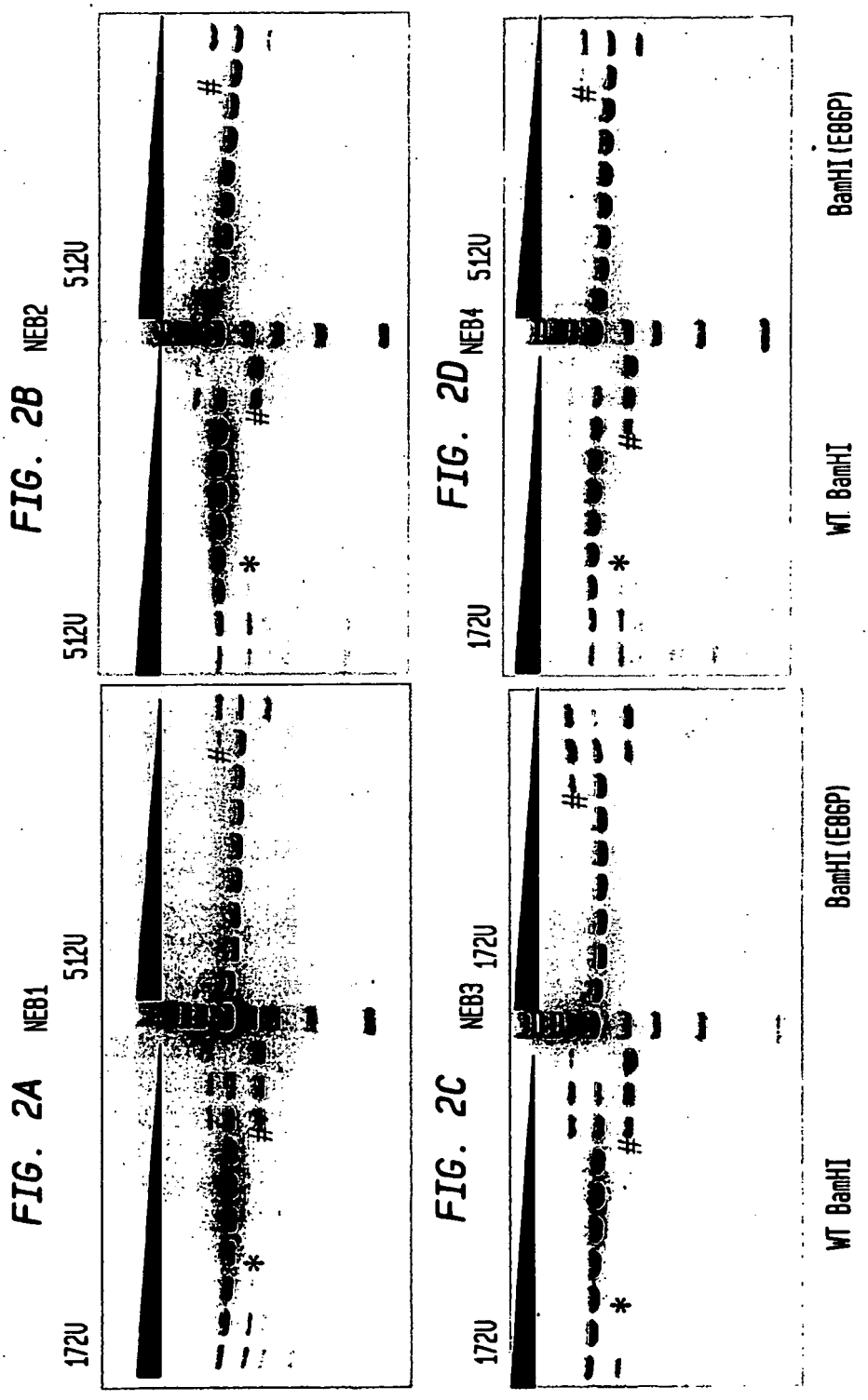
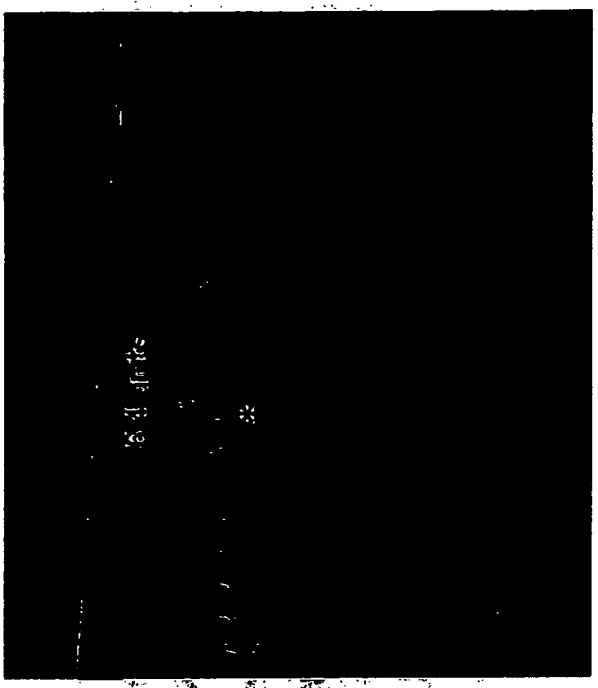


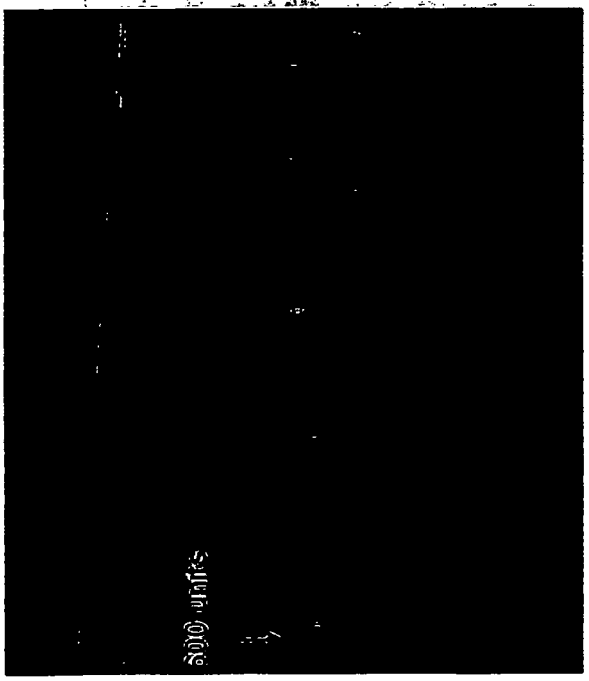
FIG. 2A
NEB1
512U
172U
WT BamHI
FIG. 2B
NEB2
512U
512U
BamHI (E86P)
FIG. 2C
NEB3
172U
172U
BamHI (E86P)
FIG. 2D
NEB4
512U
172U
WT BamHI

FIG. 3B
BamHI(E86P)



14 HOURS

FIG. 3A
BamHI(E86P)



1 HOUR

FIG. 4A

BamHI-HF(E163A/E167T)

NEB2

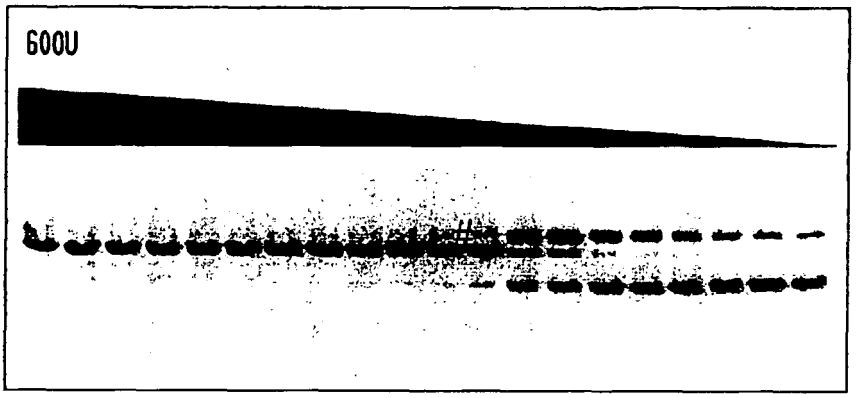
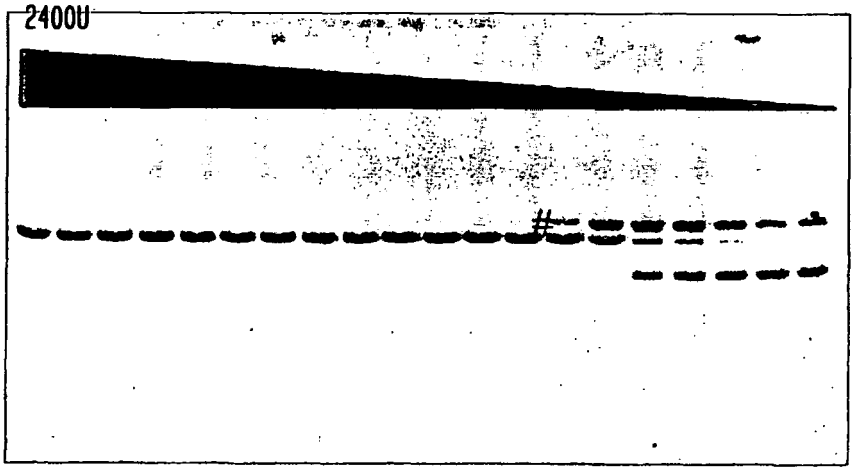


FIG. 4B

BamHI-HF(E163A/E167T)

NEB1



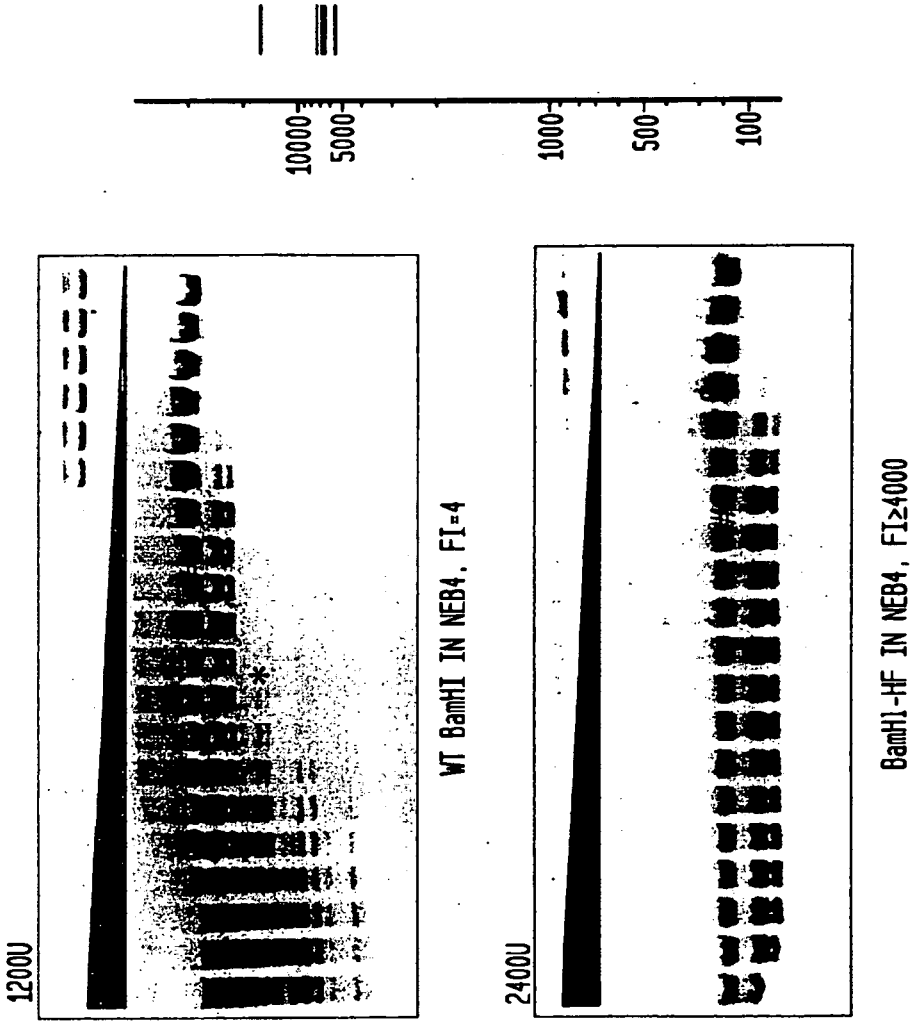


FIG. 5A
COMPARISON OF
BamHI-HF AND WT BamHI

FIG. 5B
COMPARISON OF
BamHI-HF AND WT BamHI

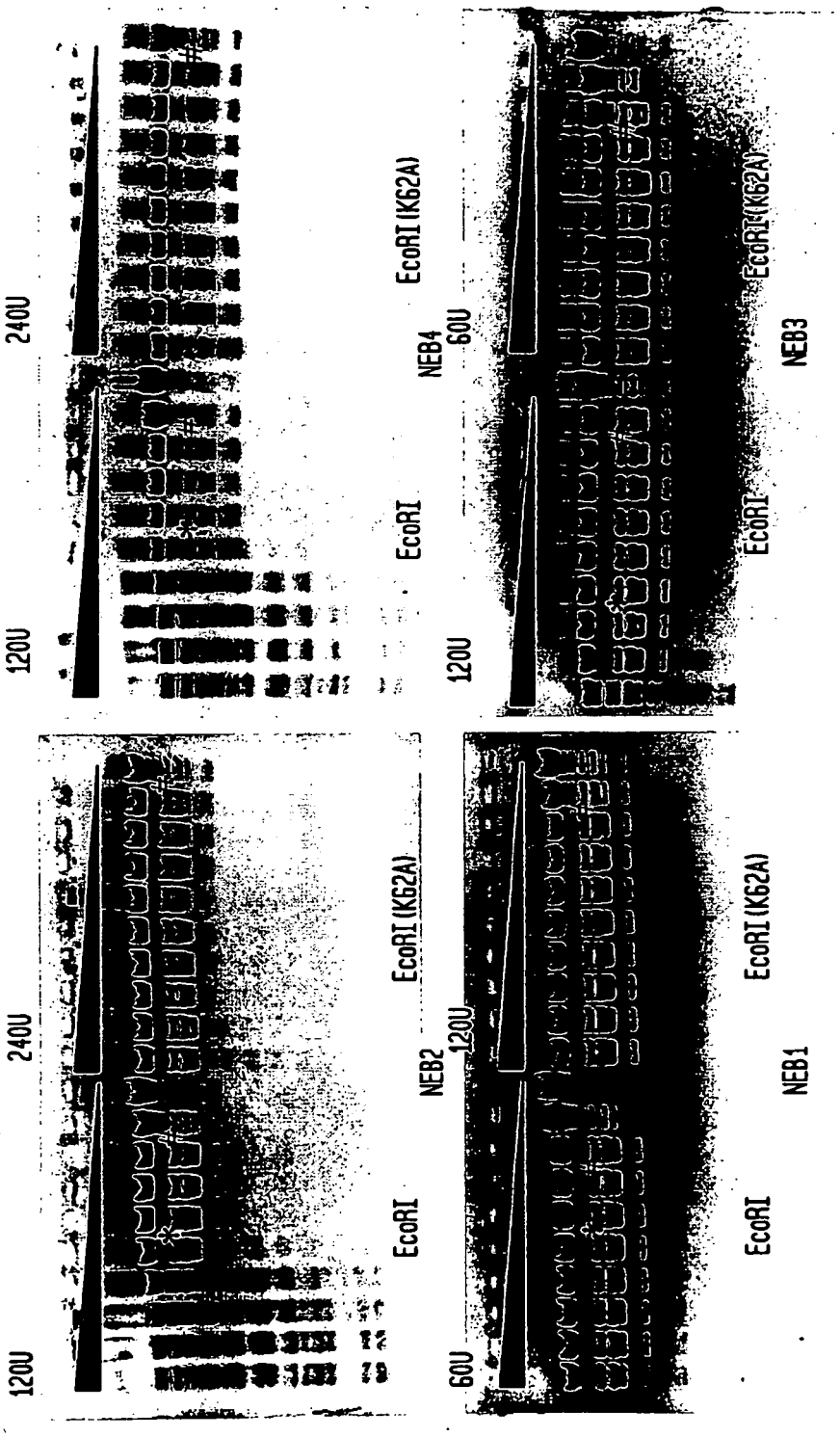


FIG. 6A
240U
120U
EcoRI
NEB2
EcoRI (K62A)
FIG. 6B
240U
120U
EcoRI
NEB4
EcoRI (K62A)
FIG. 6C
60U
120U
EcoRI
NEB1
EcoRI (K62A)
FIG. 6D
60U
120U
EcoRI
NEB3
EcoRI (K62A)

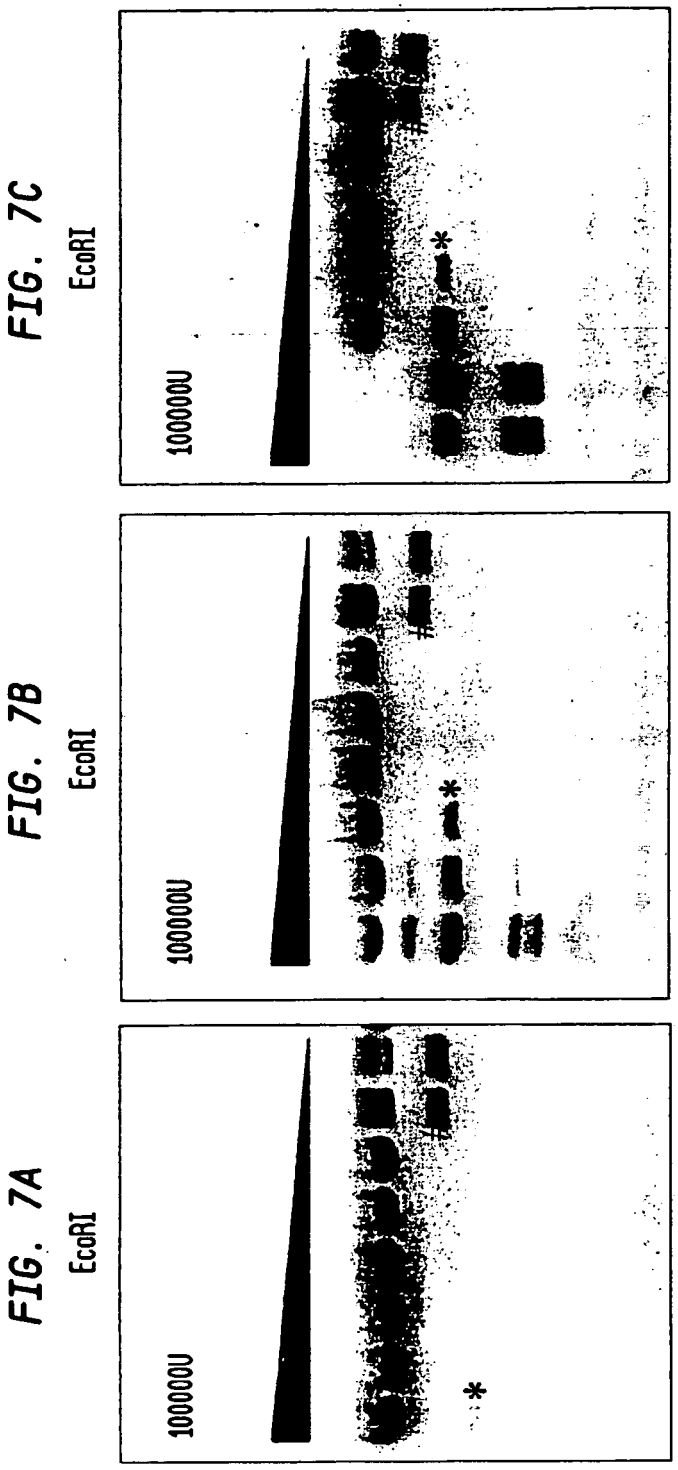


FIG. 7A
EcoRI
100000U
K62E IN NEB4
FIG. 7B
EcoRI
100000U
K62A IN NEB4
FIG. 7C
EcoRI
100000U
WT IN EcoRI BUFFER

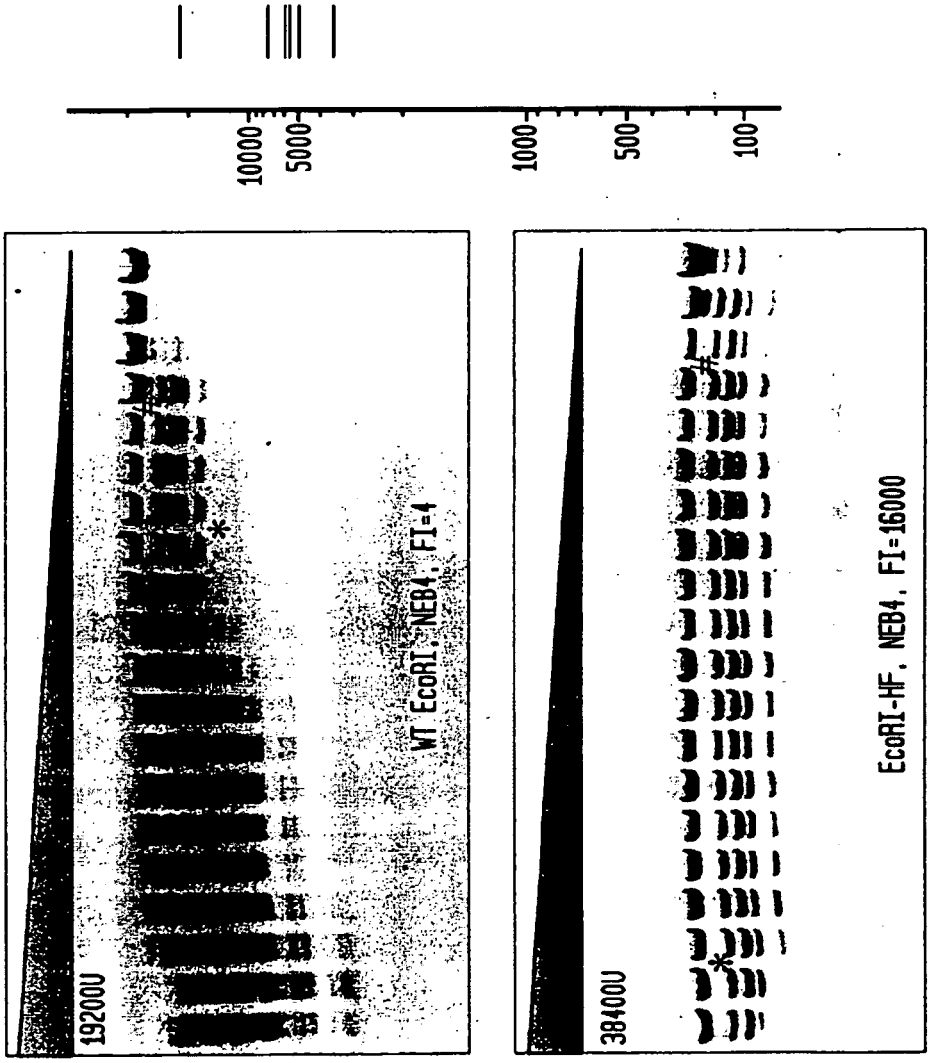


FIG. 8A
COMPARISON OF
EcoRI-HF AND WT EcoRI

FIG. 8B
COMPARISON OF
EcoRI-HF AND WT EcoRI

FIG. 9A

ScaI COMBINATION OF MUTATIONS

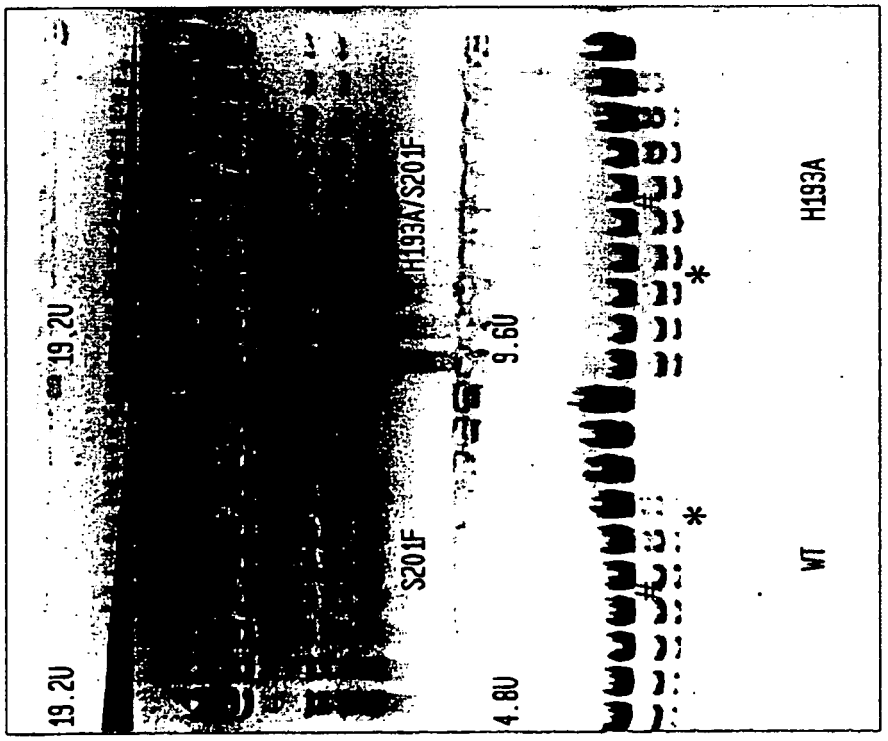
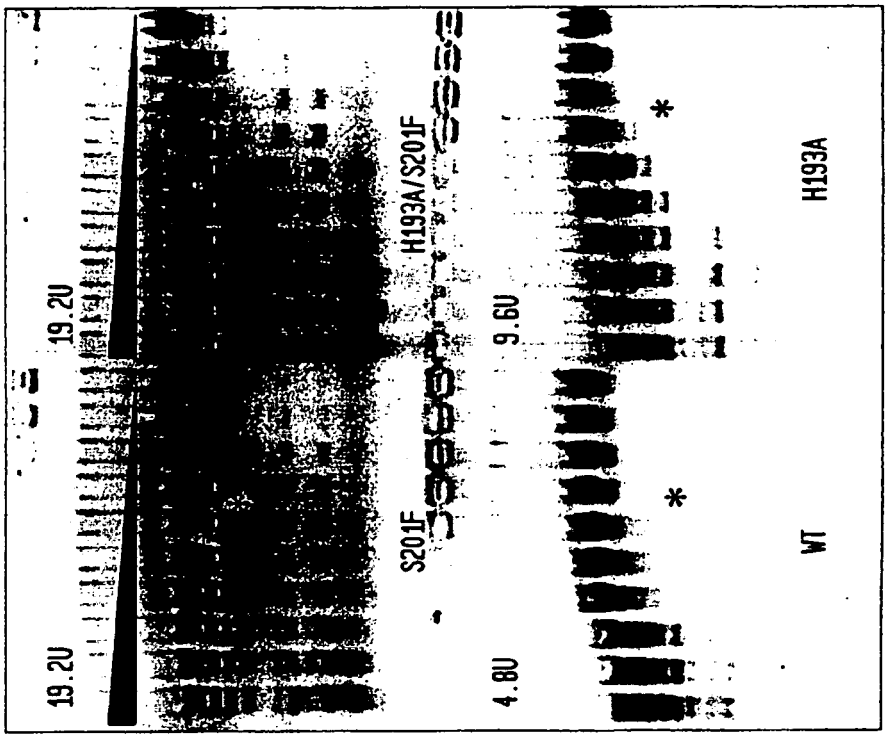


FIG. 9B

ScaI COMBINATION OF MUTATIONS



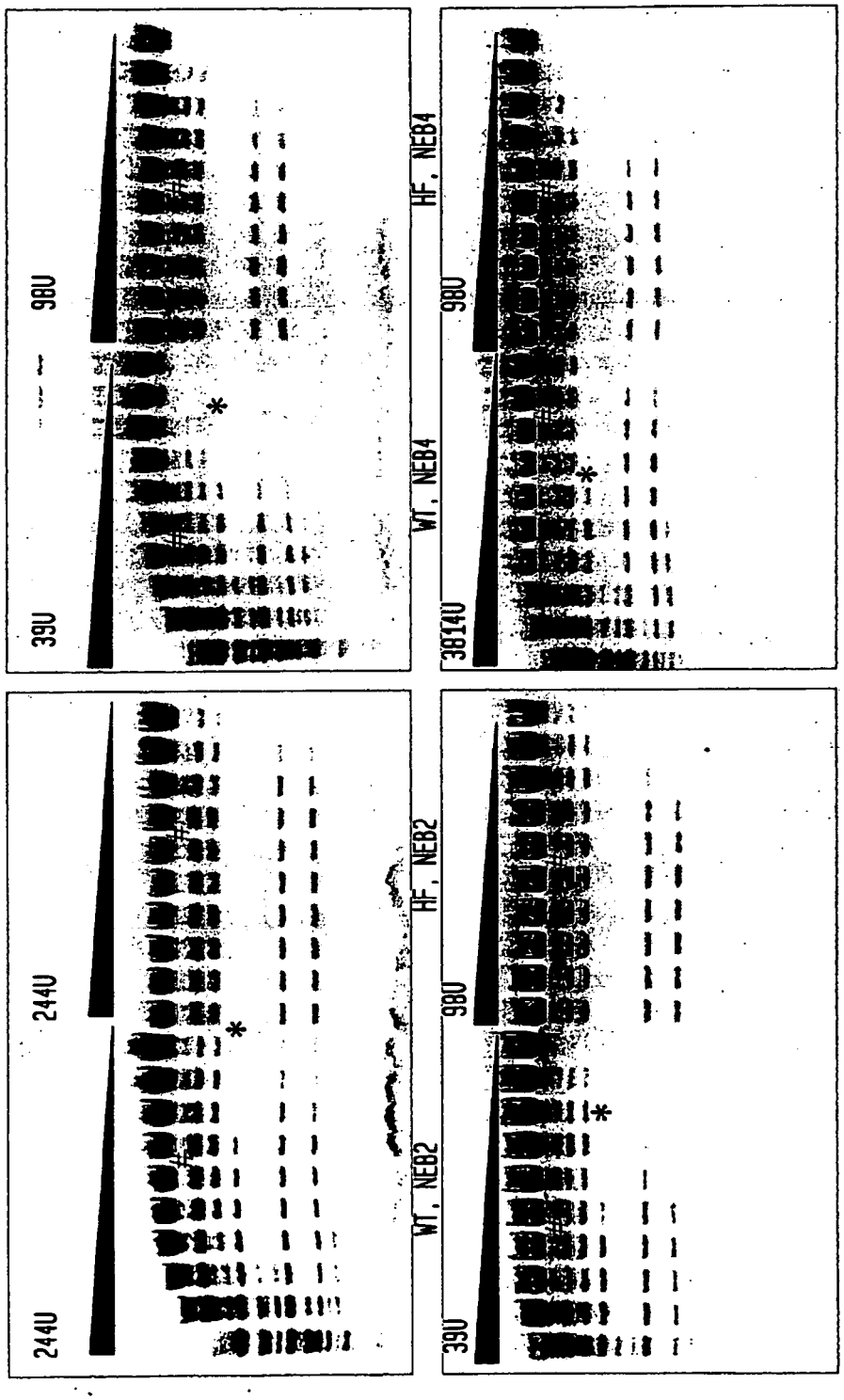


FIG. 10A
ScaI COMPARISON OF ScaI-HF AND WT ScaI

FIG. 10B
ScaI COMPARISON OF ScaI-HF AND WT ScaI

FIG. 10C
ScaI COMPARISON OF ScaI-HF AND WT ScaI

FIG. 10D
ScaI COMPARISON OF ScaI-HF AND WT ScaI

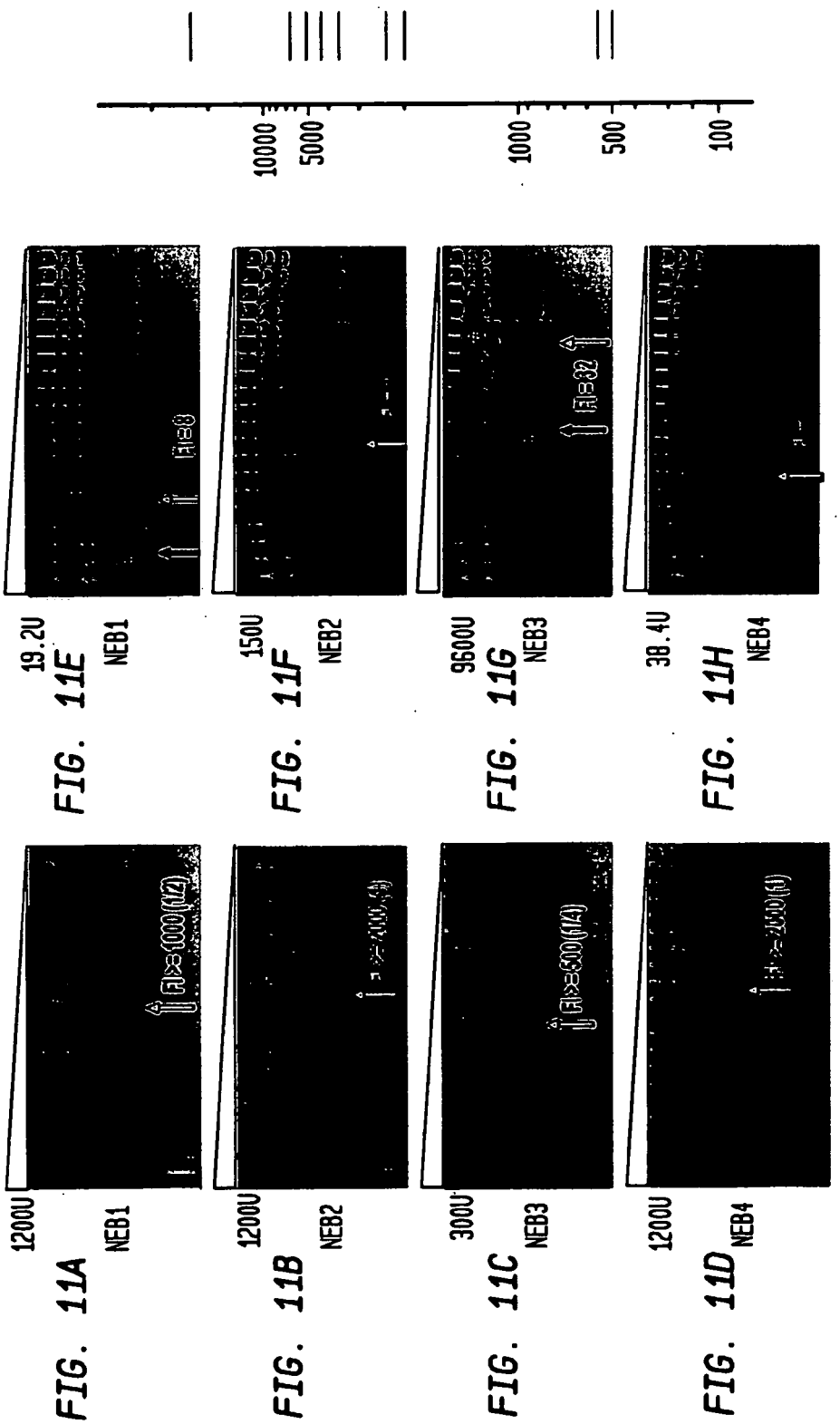


FIG. 11E FIG. 11F FIG. 11G FIG. 11H

FIG. 11A FIG. 11B FIG. 11C FIG. 11D

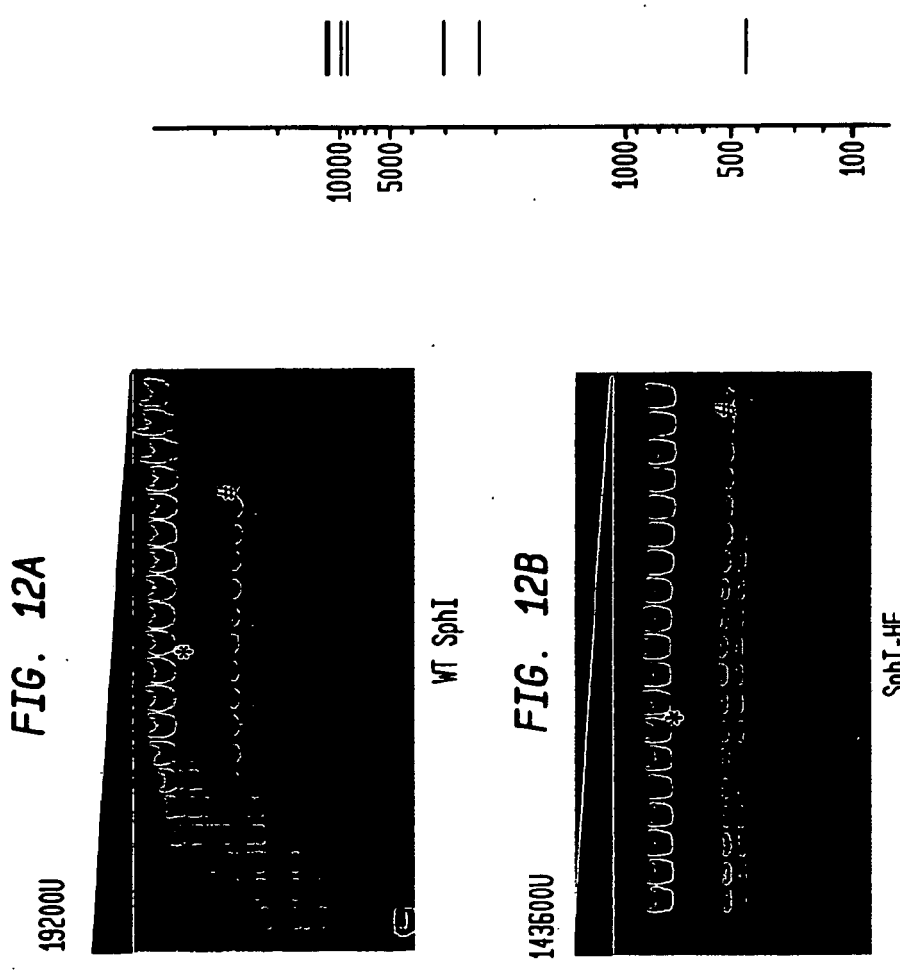


FIG. 12A

FIG. 12B

FIG. 13A

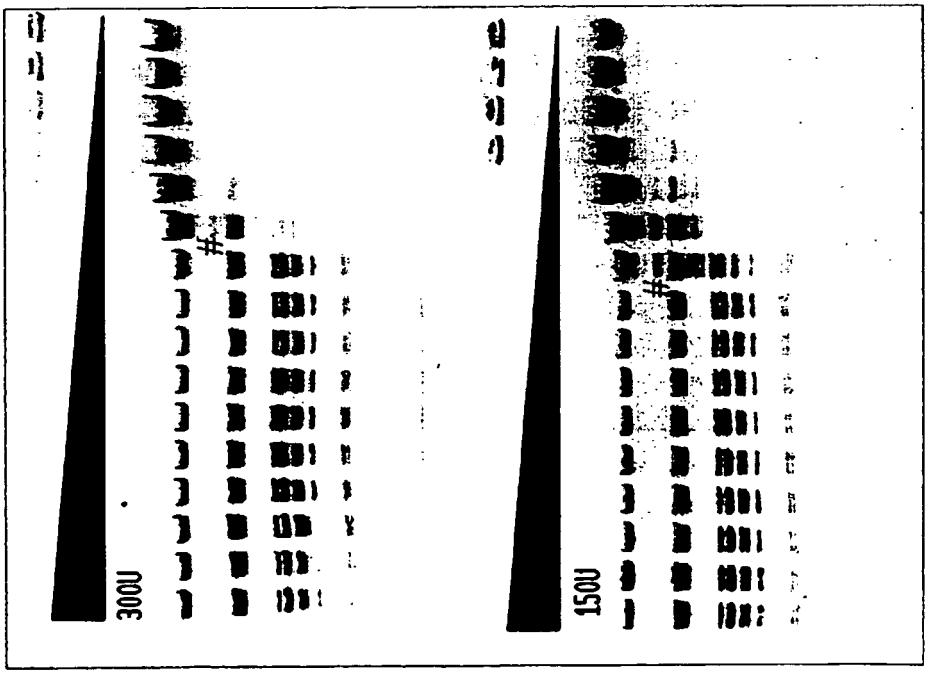
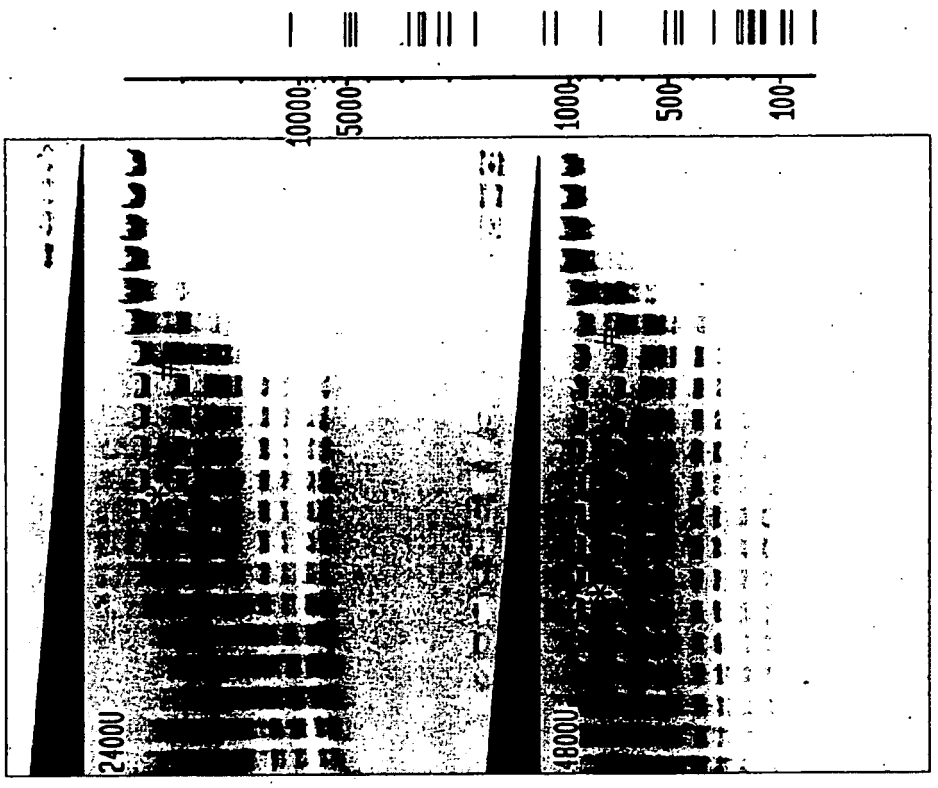


FIG. 13B



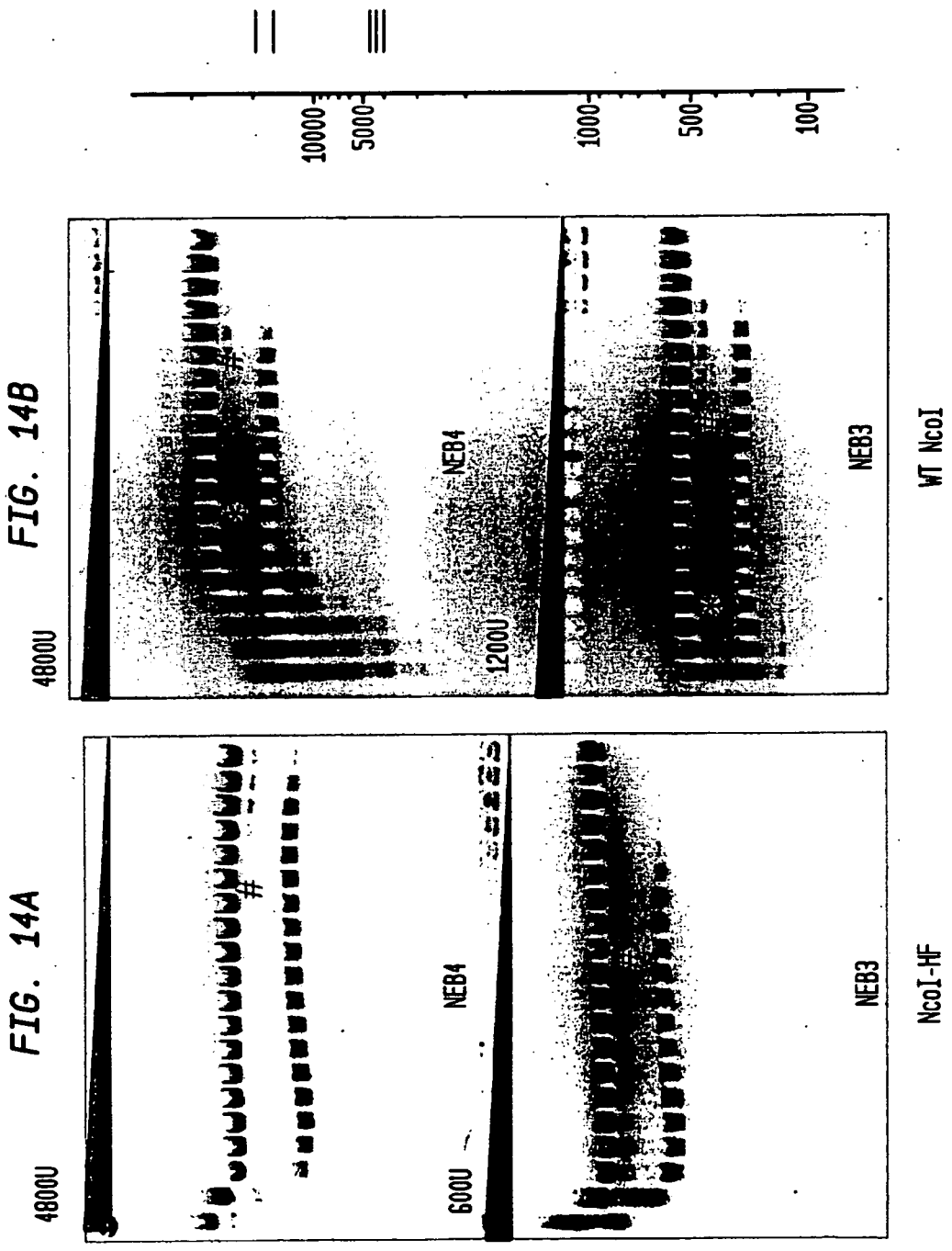

10000
5000
1000
500
100
4800U
NEB4
1200U
NEB3
WT NcoI
FIG. 14B
4800U
NEB4
600U
NEB3
NcoI-HF
FIG. 14A

FIG. 15A

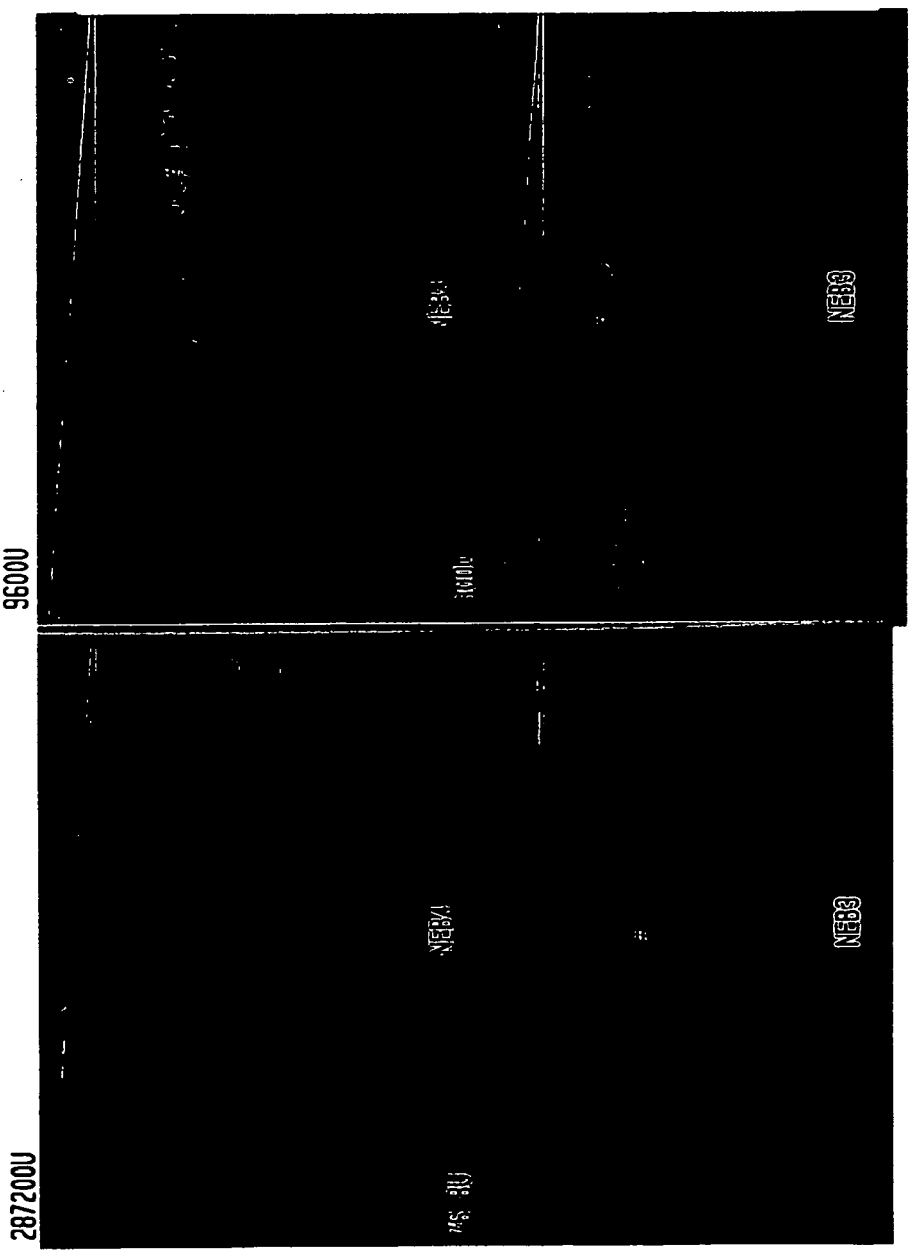


FIG. 15B

FIG. 16A

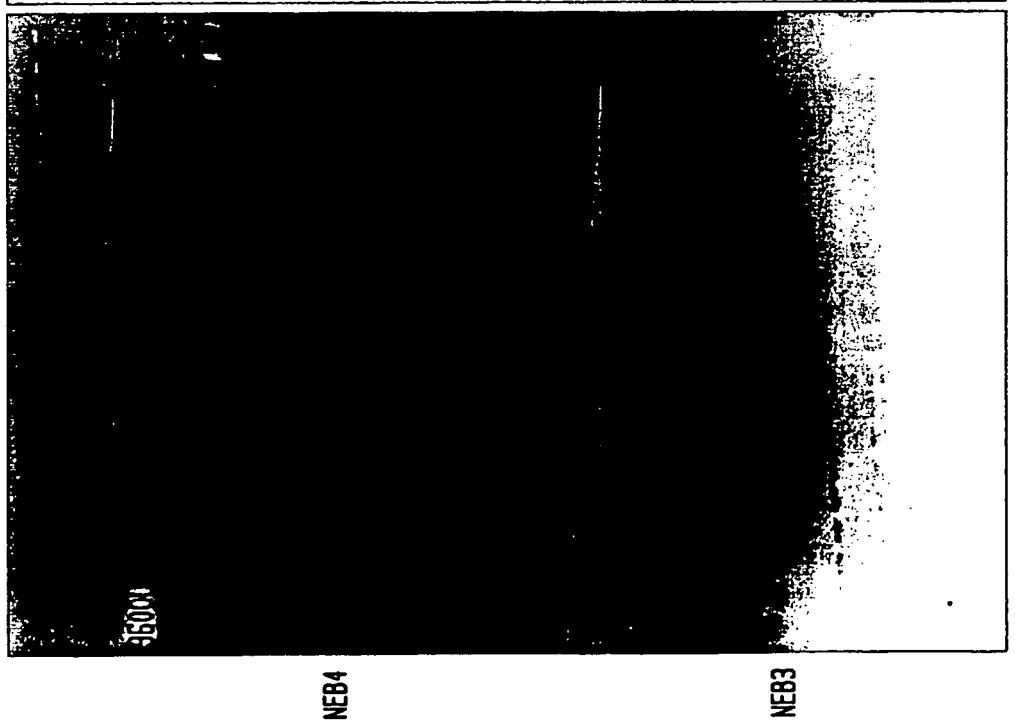


FIG. 16B

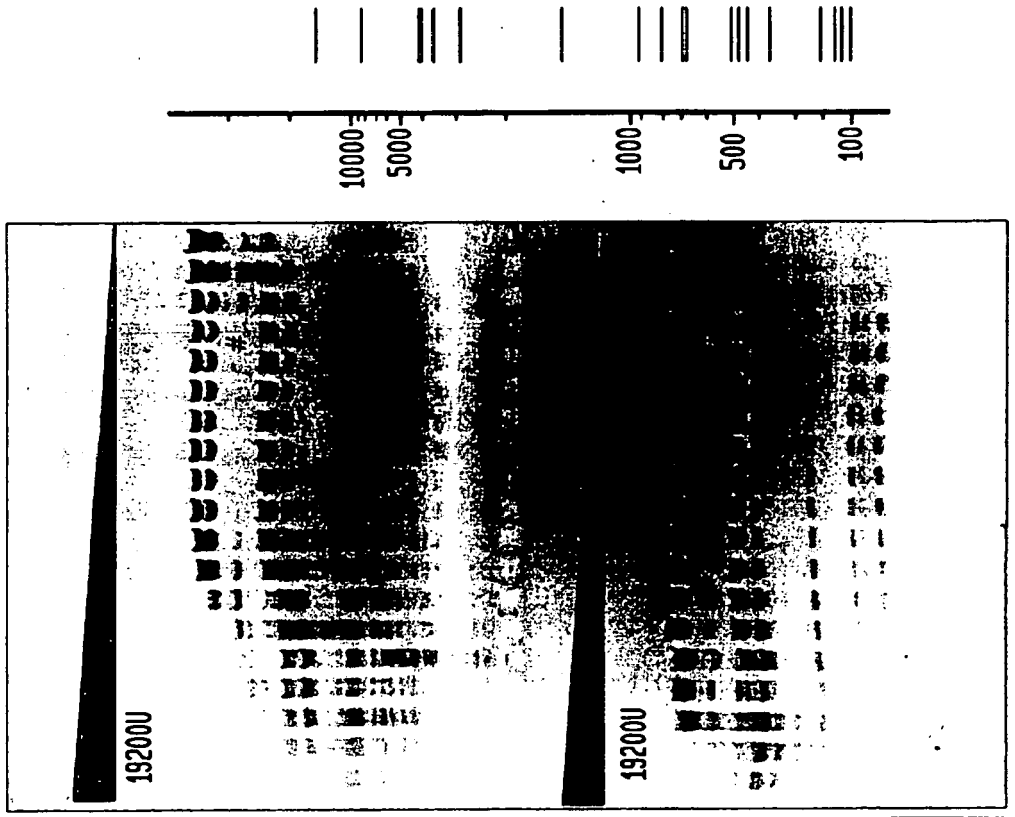


FIG. 17A

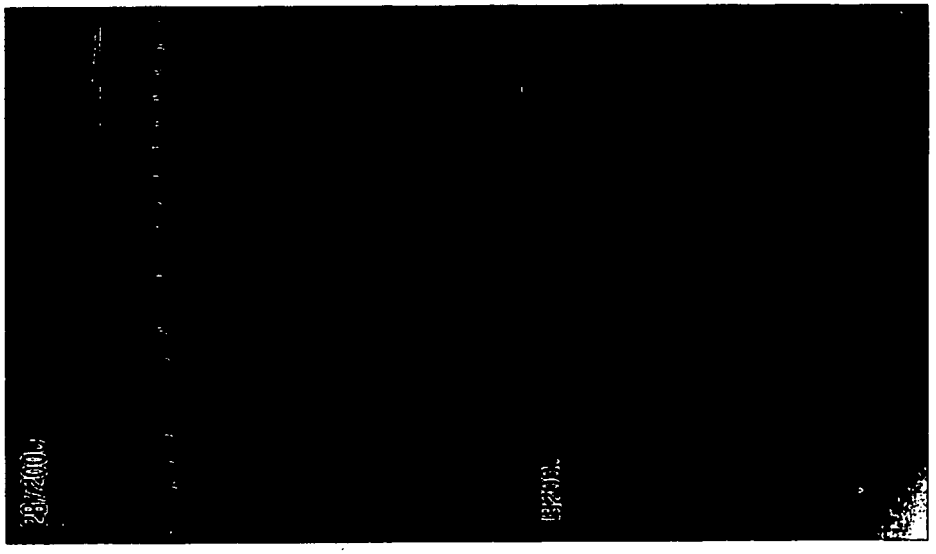


FIG. 17B

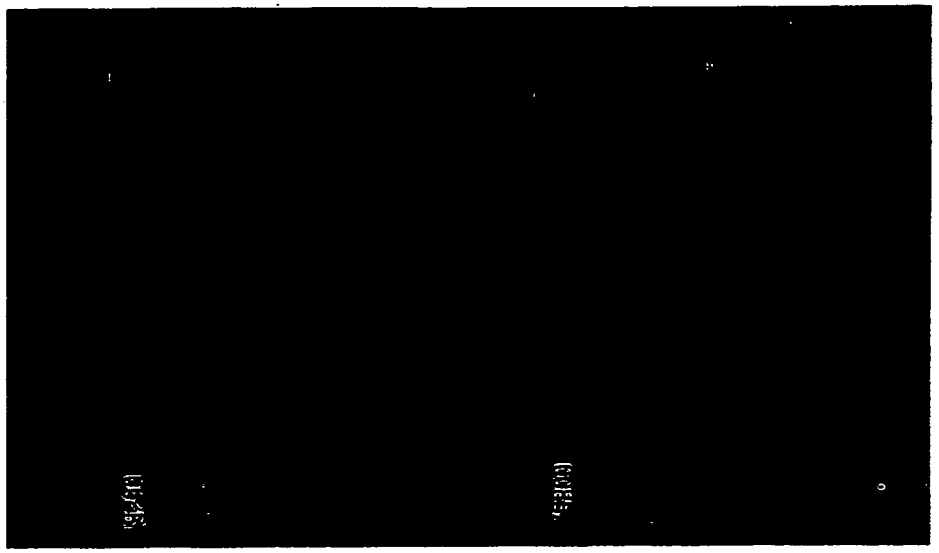


FIG. 18A

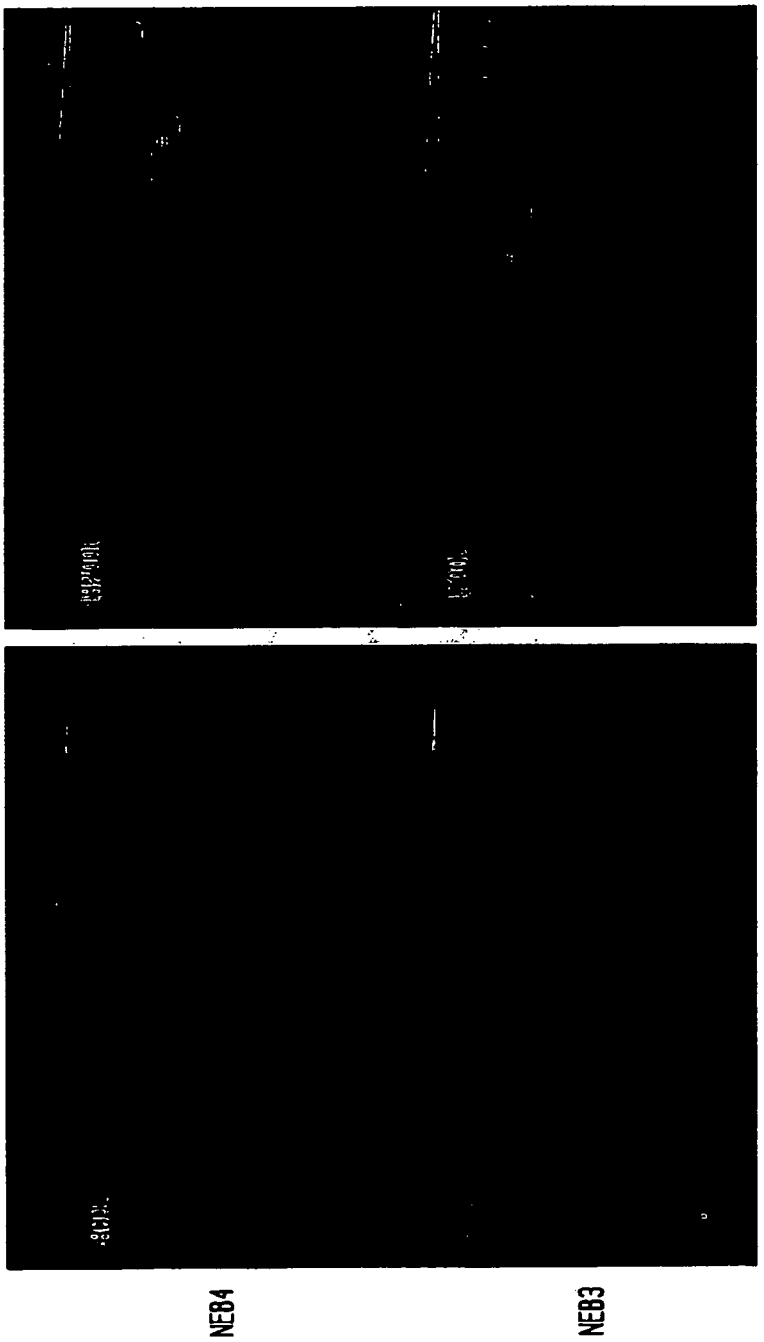


FIG. 18B

FIG. 19B

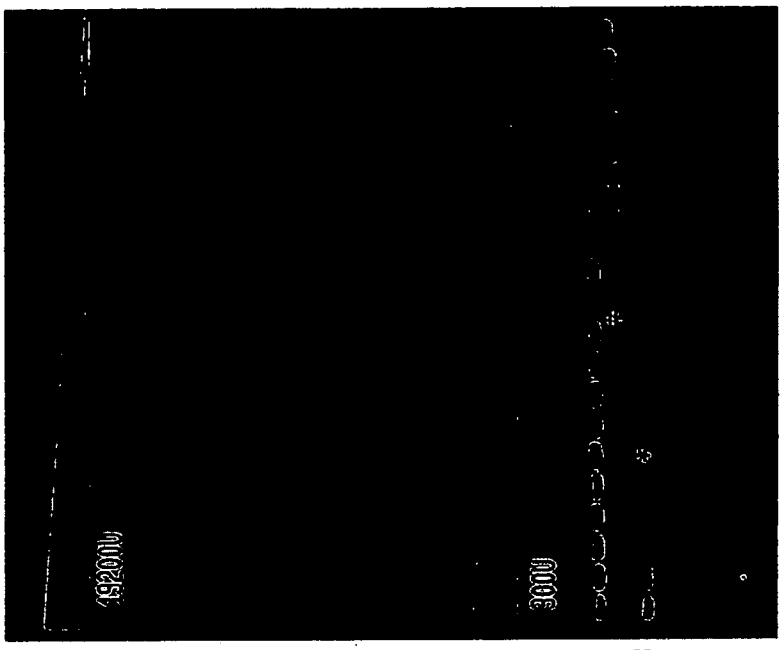


FIG. 19A

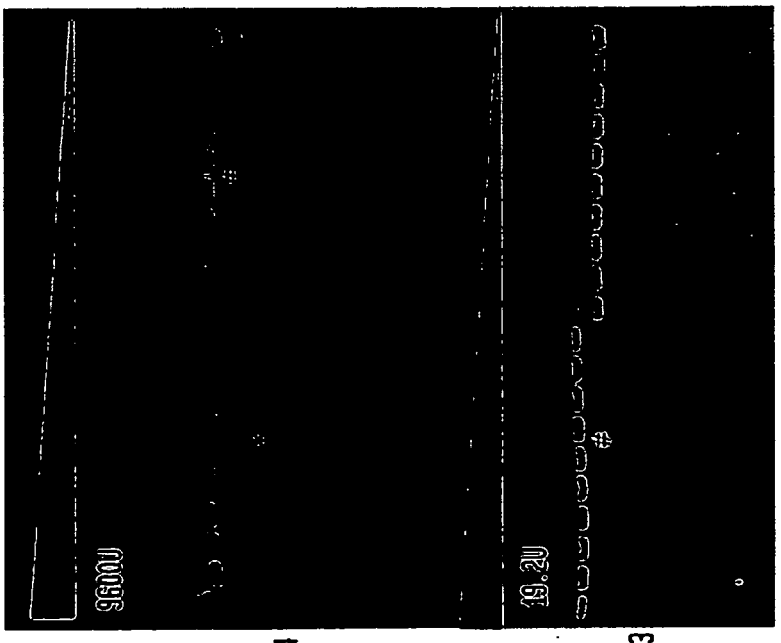


FIG. 20A

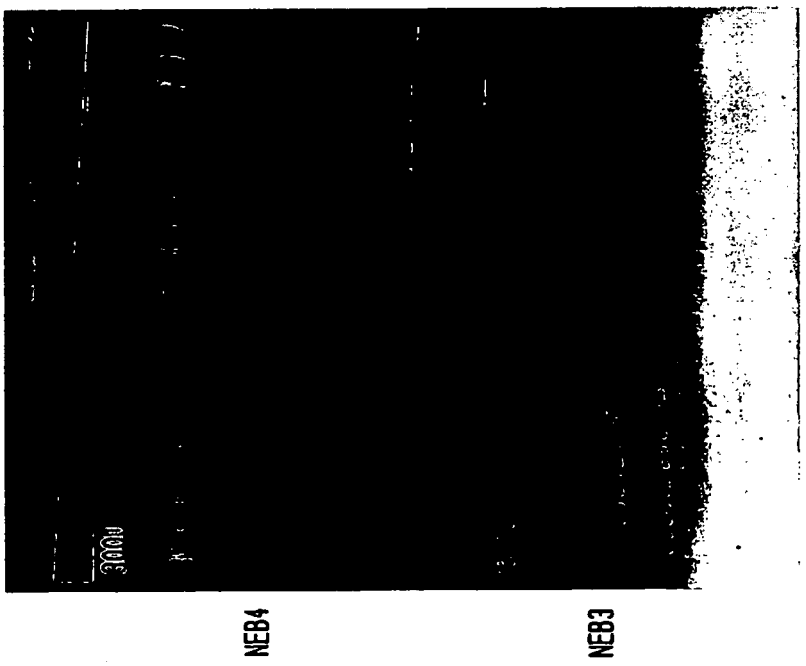


FIG. 20B

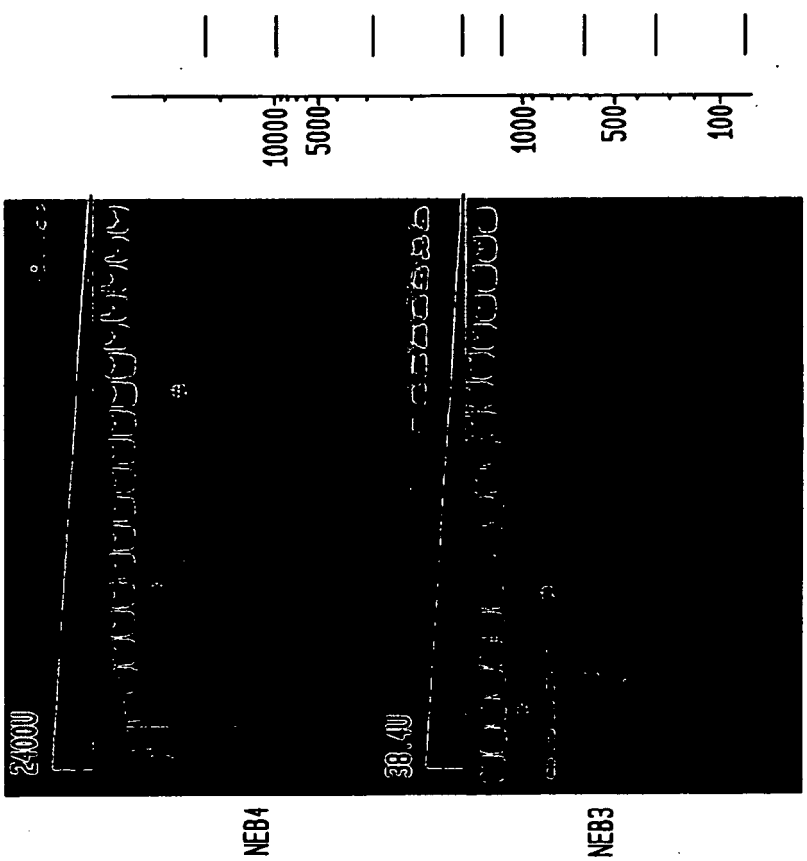


FIG. 21A

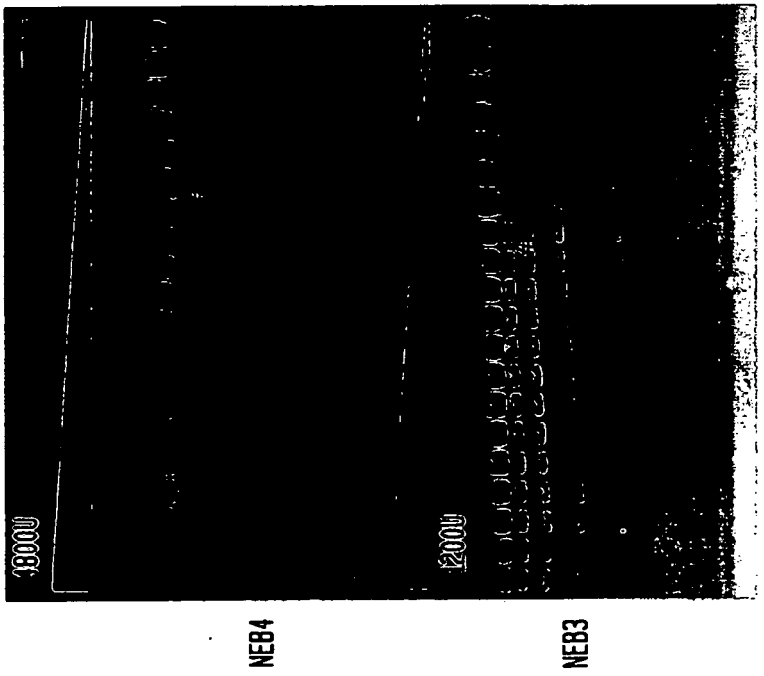
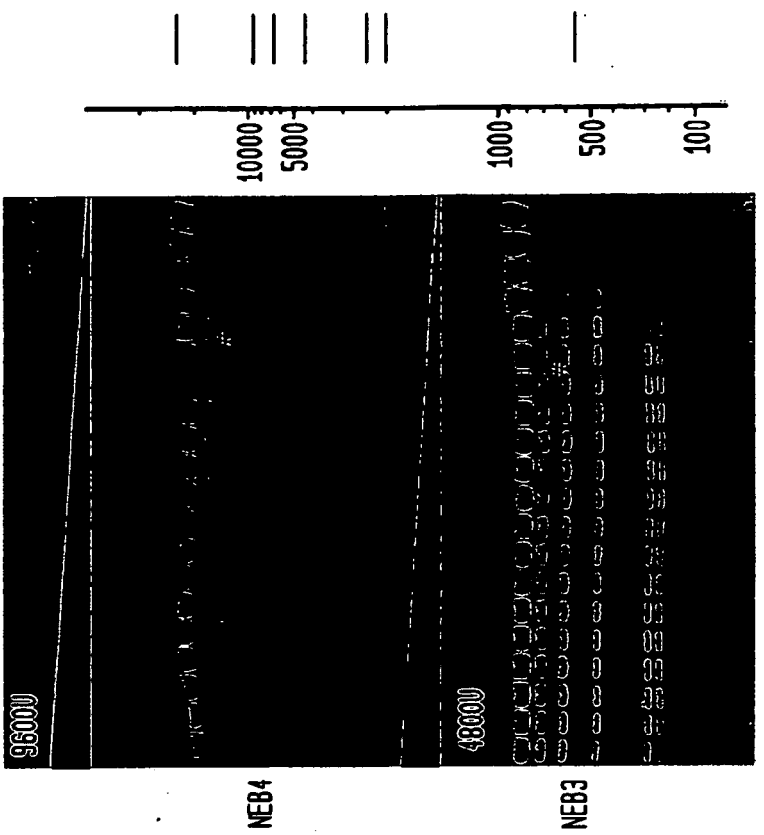


FIG. 21B



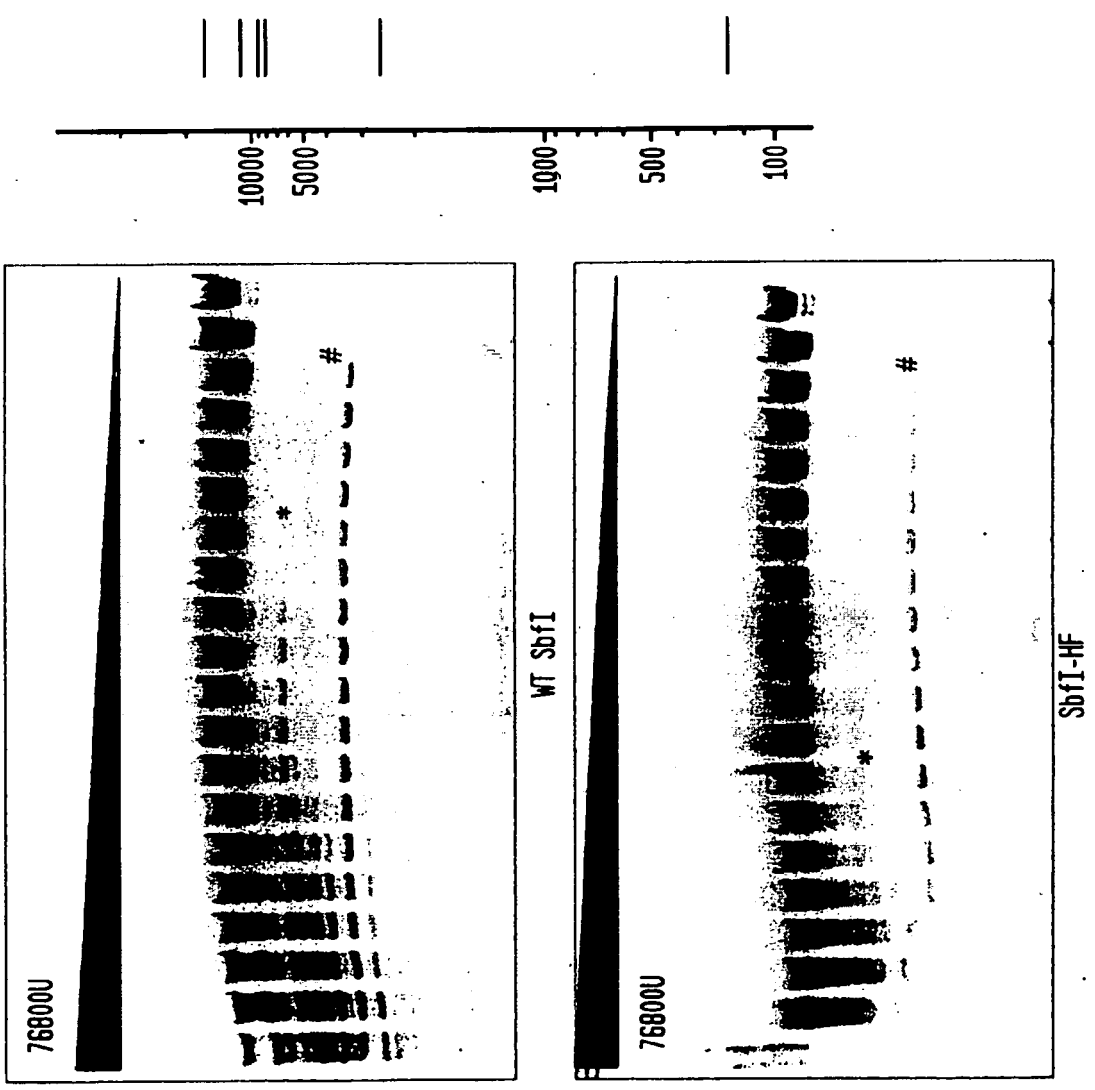


FIG. 22A

FIG. 22B

FIG. 23A

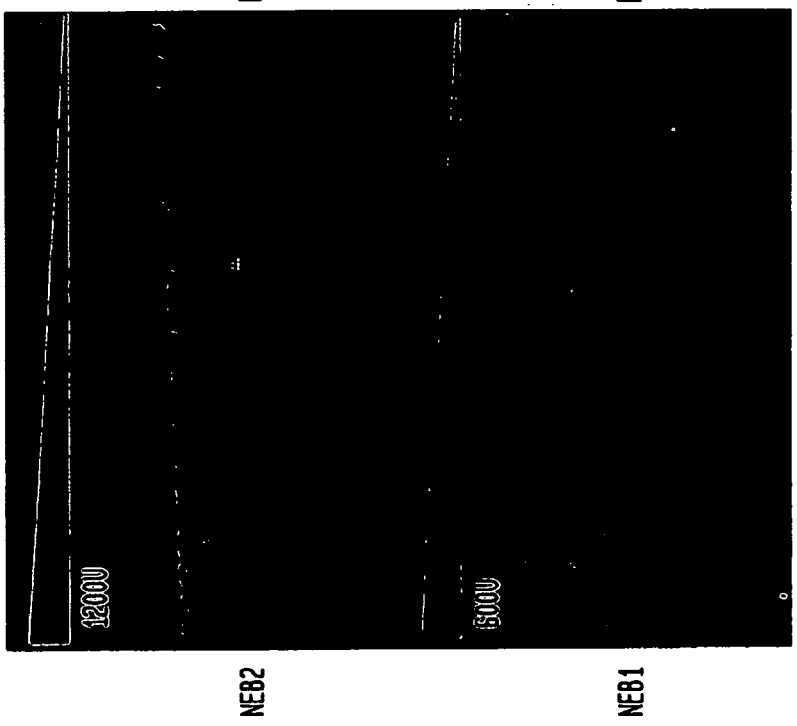
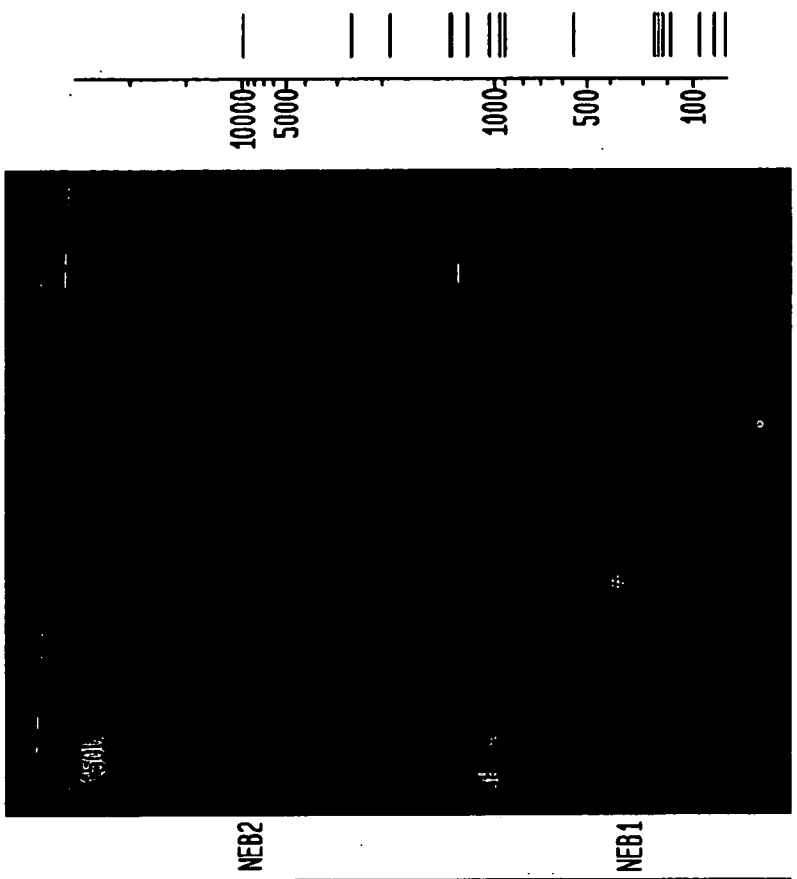


FIG. 23B



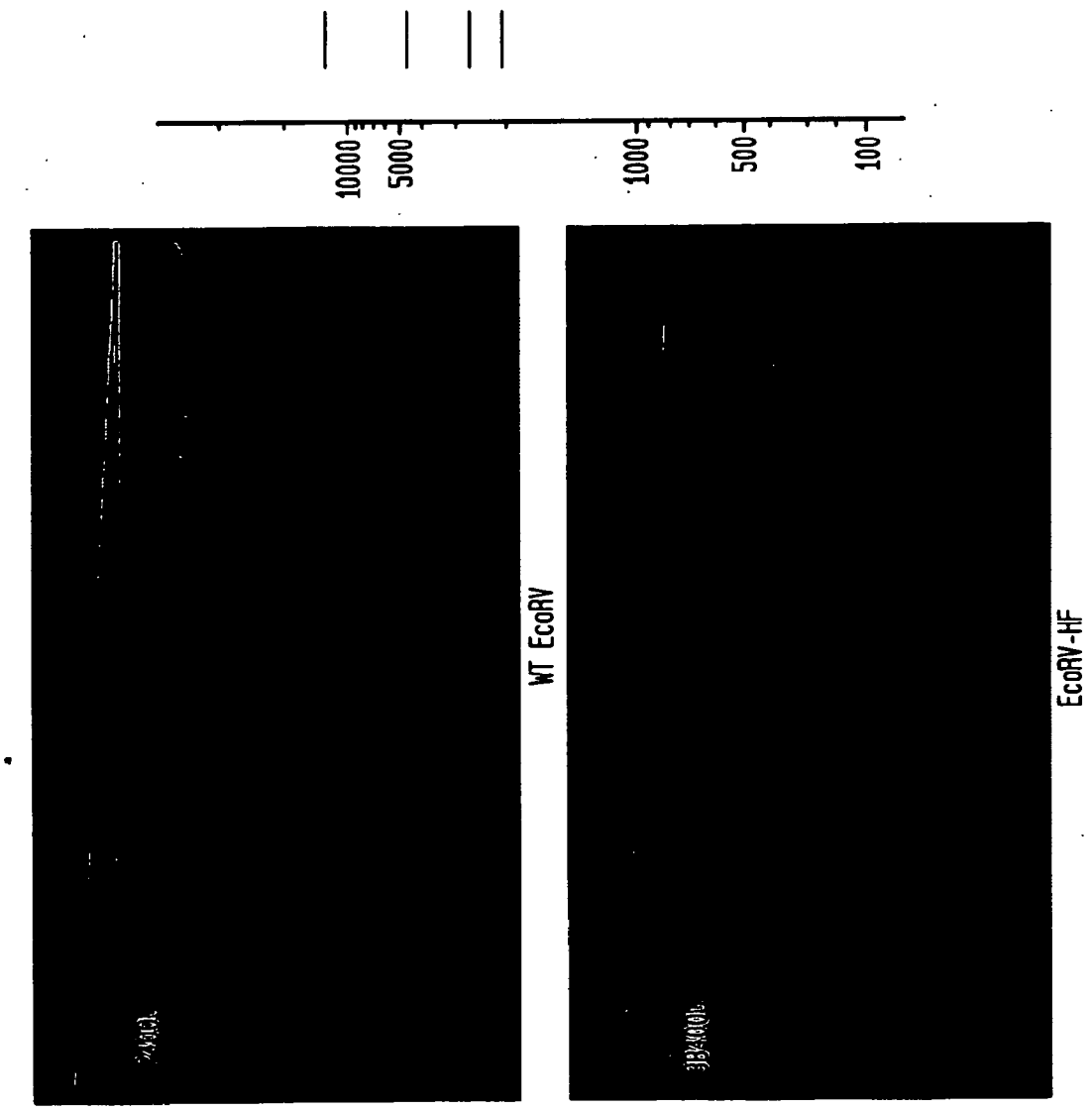


FIG. 24A

FIG. 24B

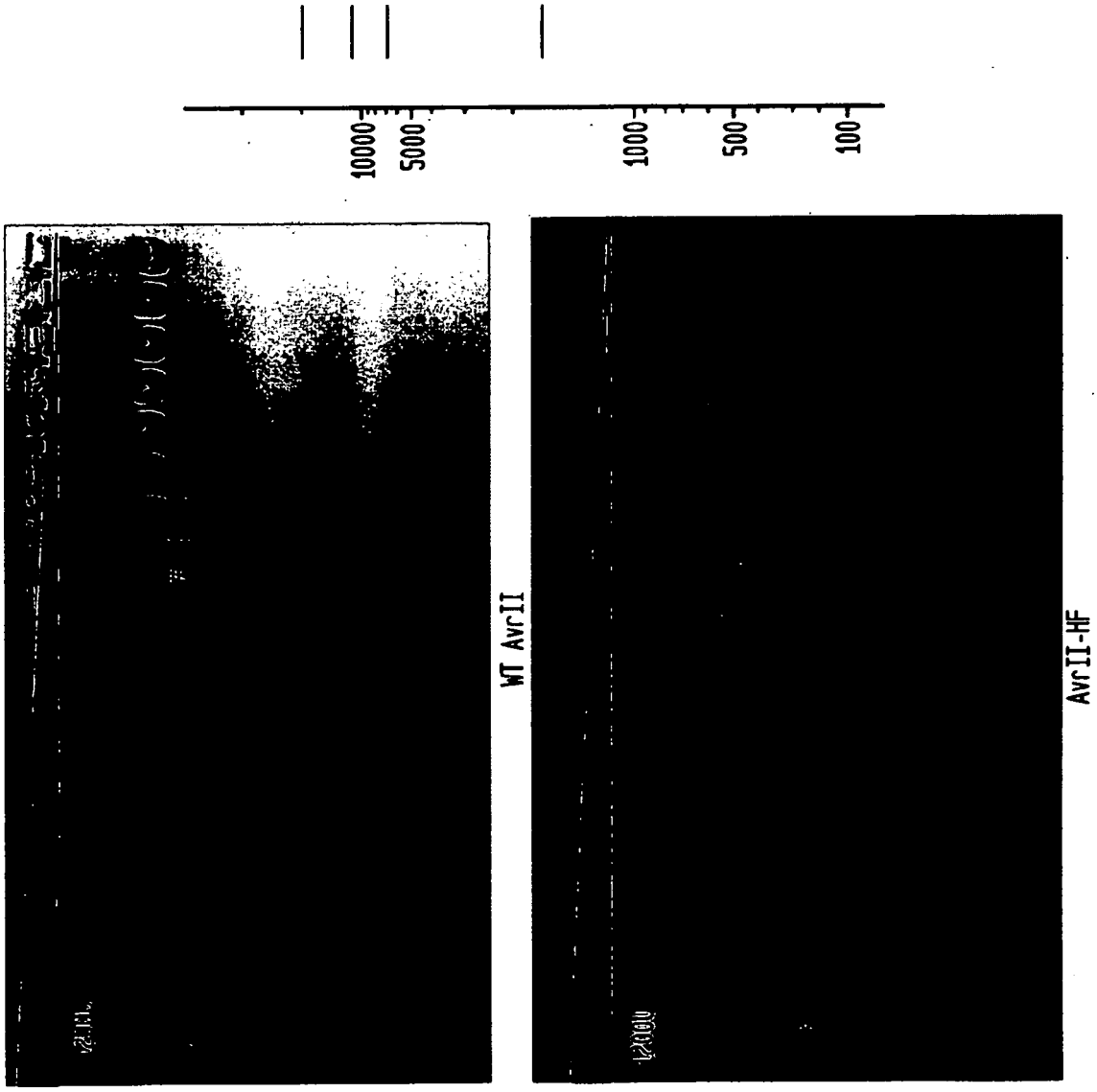


FIG. 25A

FIG. 25B

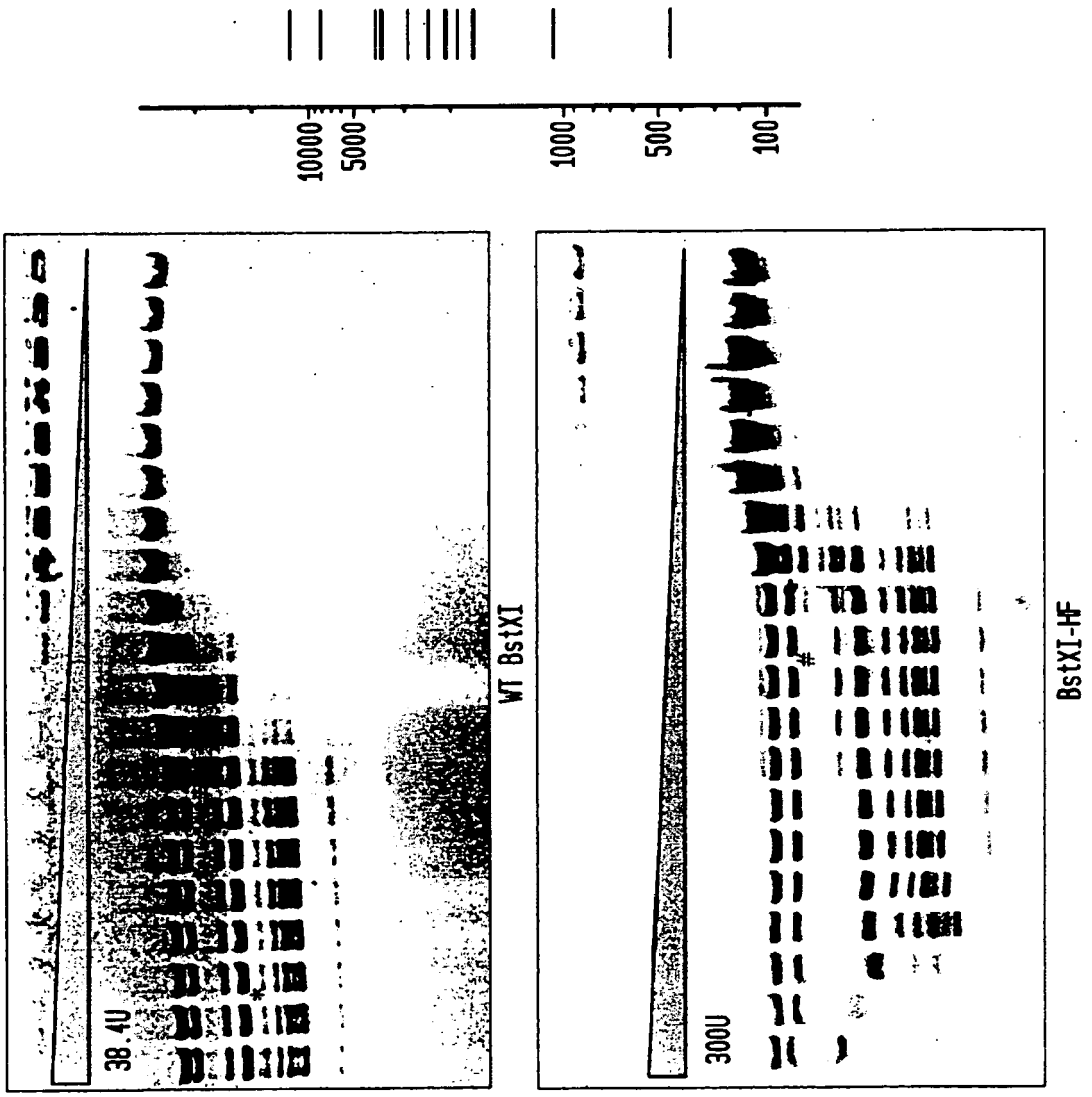


FIG. 26A

FIG. 26B

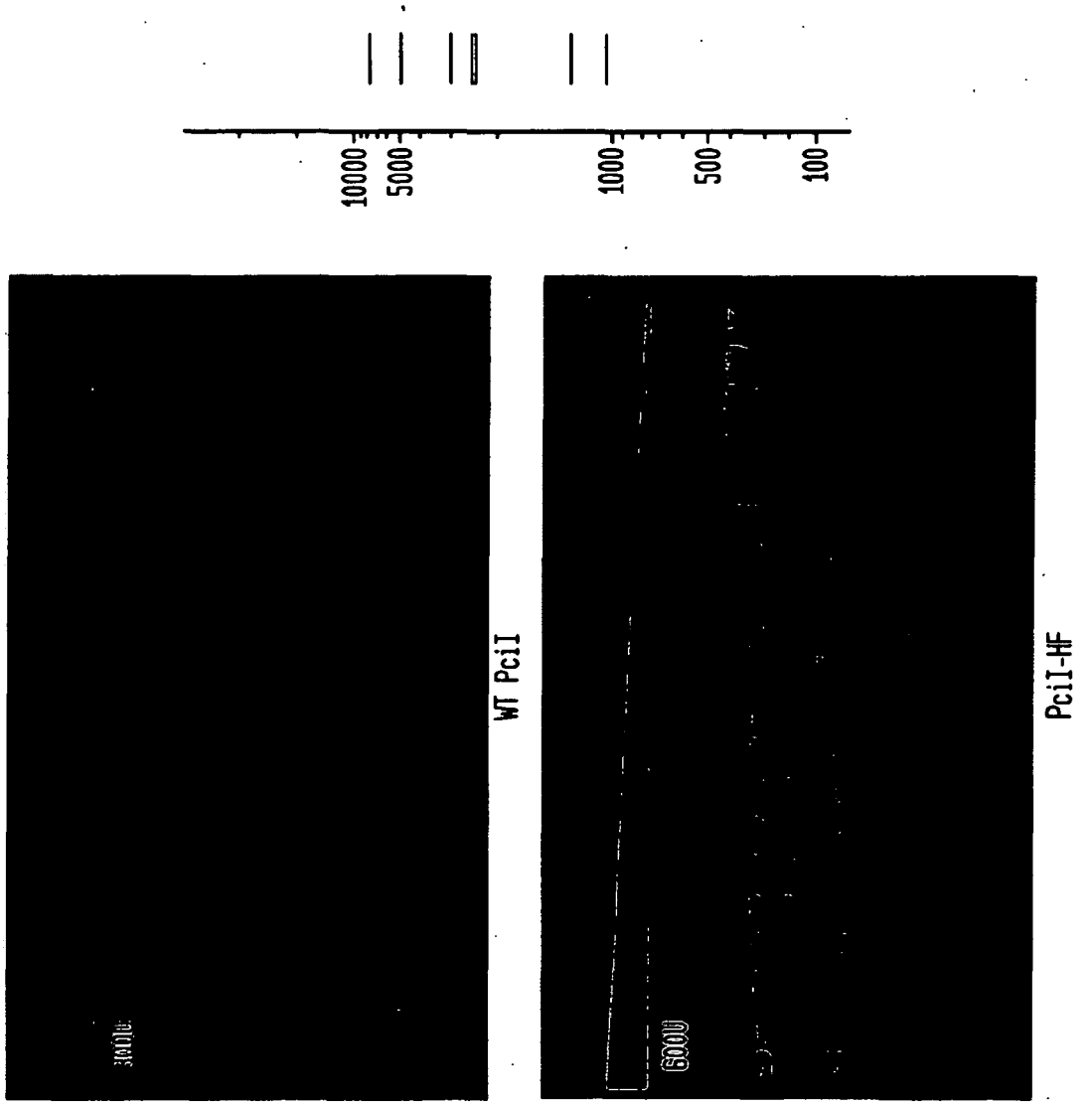


FIG. 27A

FIG. 27B

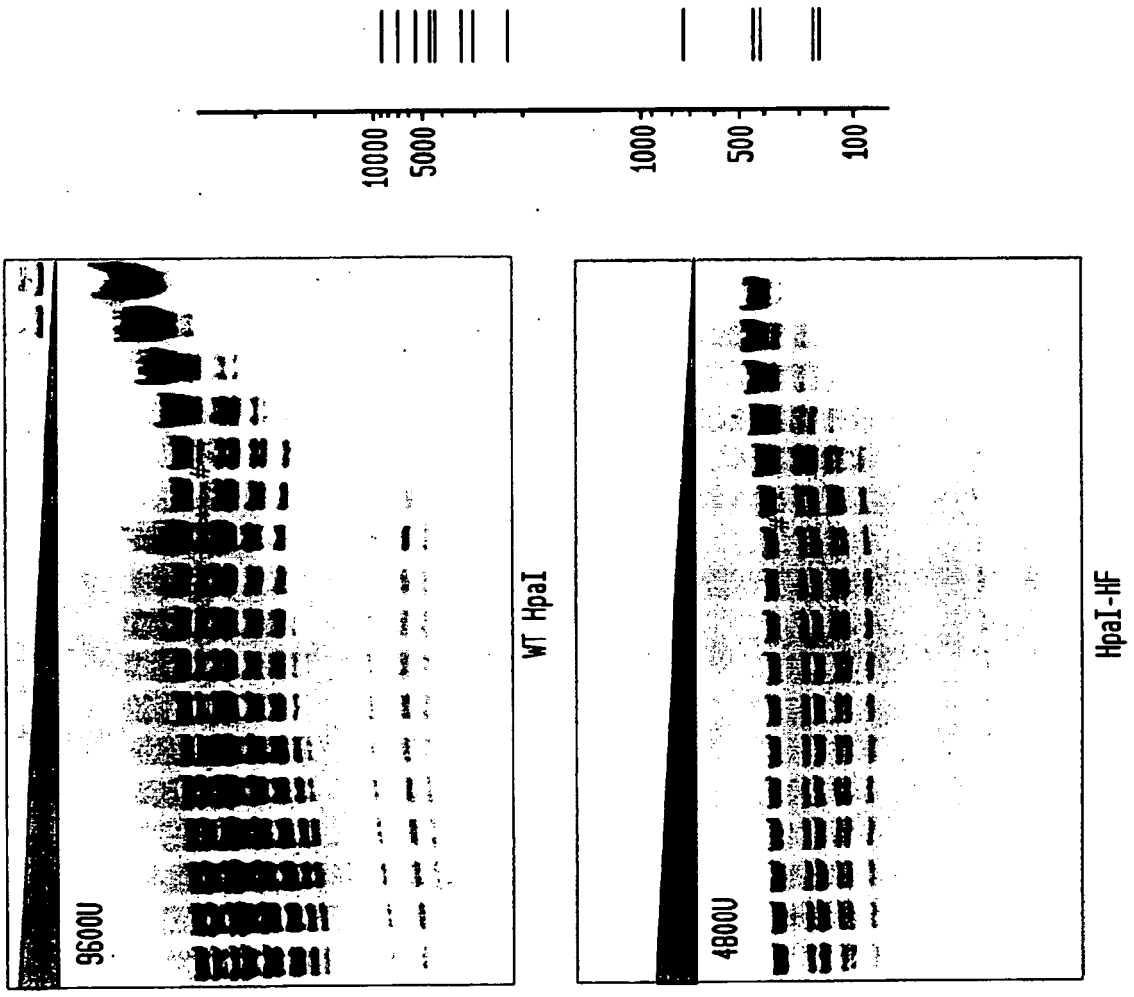


FIG. 28A

FIG. 28B

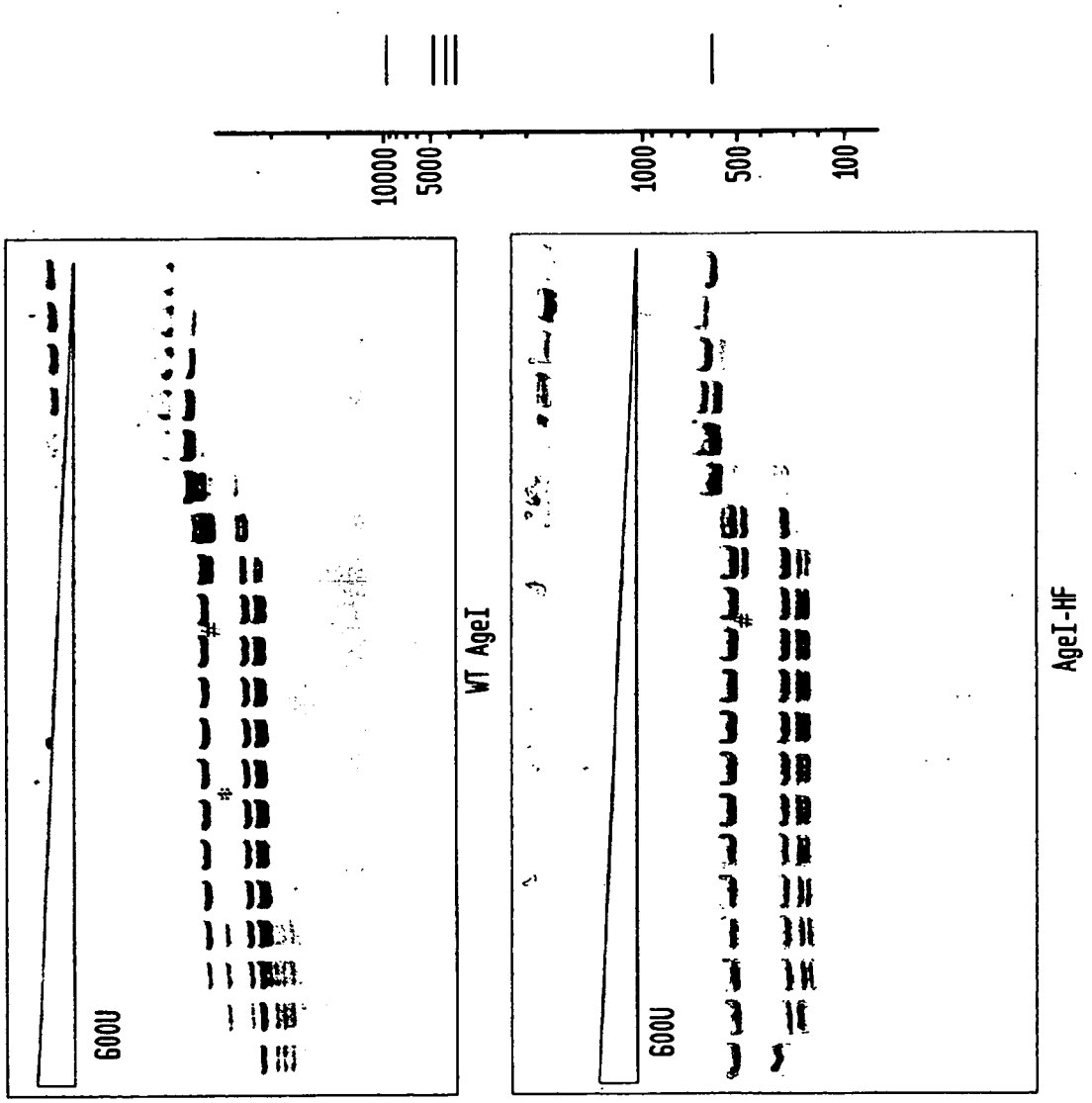


FIG. 29A

FIG. 29B

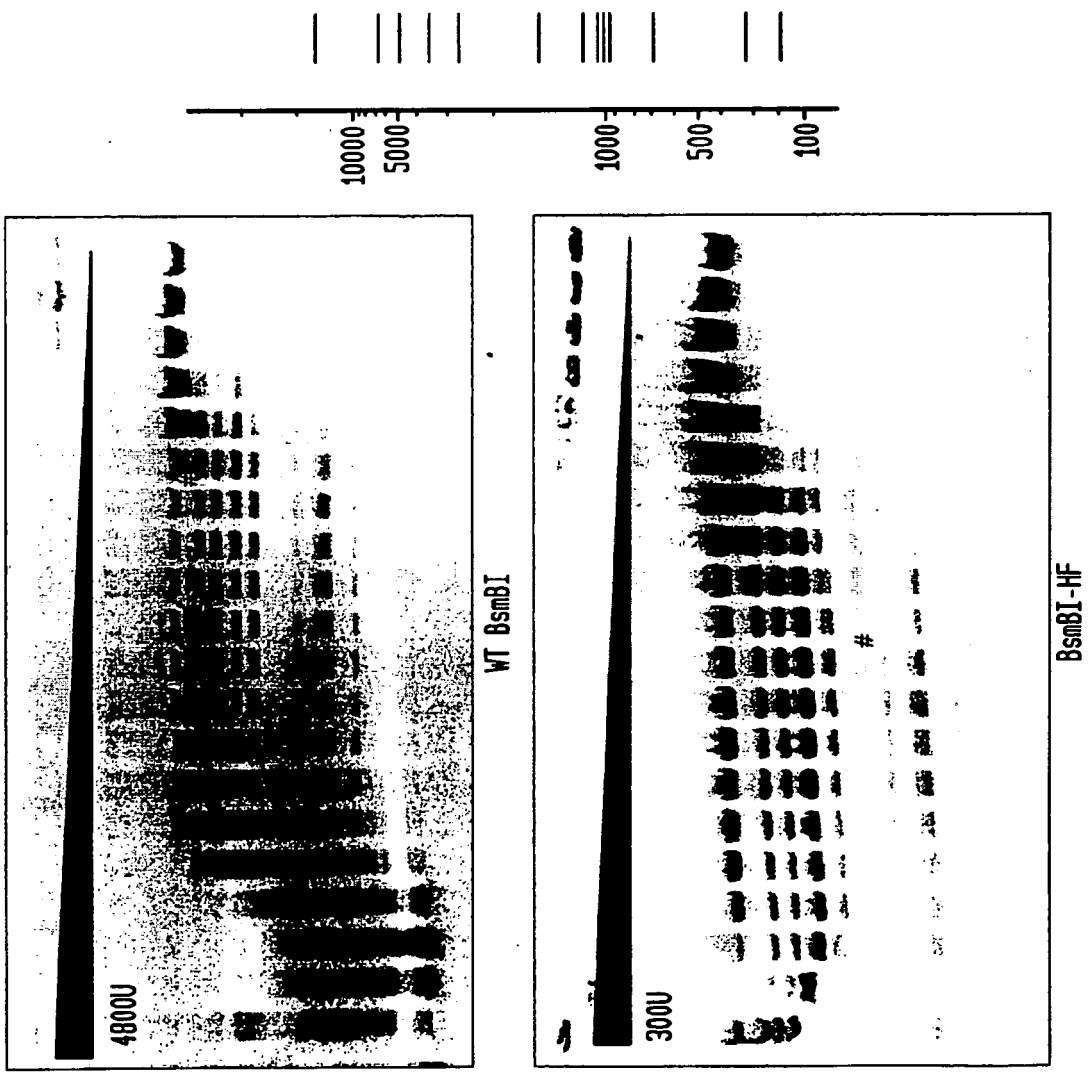


FIG. 30A

FIG. 30B

FIG. 31A

SEQ ID NO:1

```
   1  ATGATCAAGT ACTTGGGTAG CAAGCGGACG CTCGTGCCCG TCCTCGGTGA
  51  CATCGCTTCG GCCTCTGAAG CAACAGAGGC GGTTGACCTG TTCACTGGCA
 101  CGACGCGTGT GGCGCAAGAG TTCAAGCGTC GCGGGCTTCG AGTTCTTGCT
 151  AACGACATAG CGACGTACTC CGAGGTTTTA GCCCAGTGCT ATATCGCCAC
 201  CAACGGCCAG GAAGTTGACC GCCGTGCGCT CGAGGCCGCT CTGGCGGAGC
 251  TGAACGCCTT GCCCGGCGAA CCTGGATACT TCACGGAAAC CTTCTGTGAG
 301  GCTTCTCGCT ACTTCCAGCC CAAGAACGGG GCTCGGGTGG ATGCAATCAG
 351  GAATGCGATC GACGACCGGT ACGCGGACTC ATGGATGCGA CCGATCCTCC
 401  TCACGAGCTT GATGCTTGCG GCCGACCGCG TCGACTCCAC TACCGGAGTG
 451  CAGATGGCTT ACCTGAAGCA GTGGGCCGCG CGTGCGCACA ATGATCTAGA
 501  GTTGCGGCTT CCAGACCTAA TCGCAGGTGA CGGTGACGCT GCTCGTGAGG
 551  ATGCGGTGAC TCTCGCACAA GAGCTGCCTC GCGTCCAGCT GATGTACCTT
 601  GATCCTCCCT ATAACCAGCA CAGGTACTTC ACCAACTACC ATATTTGGGA
 651  GACCCTGATT CGTTGGGATG CCCCTGAGAG TTATGGGATC GCCTGTAAGC
 701  GCATTGACTC TCGAGATGAT GCCACCAAGA GCCCCTATAA TATGAAGCGG
 751  CGAATGCCCG ACGAGATGCG TCGCCTGCTG ATGACCATCA AGGCGGACCT
 801  CGCGGTTGTA TCTTACAACA ATGAGTCGTG GATTGATCCG GAGACGATGA
 851  TGTCGACCCT GCGCGATGCG GGATATGAGG ACGTGCGTCT GCTCGCTTTC
 901  GACTATAAGC GCTACGTTGG GGCTCAAATC GGGATCTACA ATCCCTCCGG
 951  GGAAAAGGTC GGTCGTGTGA GTCACCTCCG AAACATCGAG TATCTCTTTC
1001  TTGCGGGACC AACGGAGCGC GTTGAGGTGT GCGCCGCGAG TGTTGAACAC
1051  CGAGCACTAC CCAAGGAACC GGAACTCACC GCGTTCTAG
```

FIG. 31B

SEQ ID NO:2

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MIKYLGSKRT LVPVLGDIAS
ASEATEAVDL FTGTTRVAQE
FKRRGLRVLA NDIATYSEVL
AQCYIATNGQ EVDRRALEAA
LAELNALPGE PGYFTETFCE
ASRYFQPKNG ARVDAIRNAI
DDRYADSWMR PILLTSLMLA
ADRVDSTTGV QMAYLKQWAA
RAHNDLELRL PDLIAGDGDA
AREDAVTLAQ ELPRVQLMYL
DPPYNQHRYF TNYHIWETLI
RWDAPESYGI ACKRIDSRDD
ATKSPYNMKR RMPDEMRRLL
MTIKADLAVV SYNNESWIDP
ETMMSTLRDA GYEDVRLLAF
DYKRYVGAQI GIYNPSGEKV
GRVSHLRNIE YLFLAGPTER
VEVCAASVEH RALPKEPELT
AF
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FIG. 32
SEQ ID NO:3

TTGGAGAATT TTTTGAATAA TTTAGATATT AAAACCTTAG GGCAGGTTTT CACCCCTAAA
AAGATAGTGG ATTTCATGCT CACTCTCAAG CACAATCATG GGAGTGTTTT AGAGCCAAGC
GCGGGCGATG GGAGTTTTTT AAAGCGCTTA AAAAAGGCTG TAGGGATTGA AATCGATCCT
AAAATCTGCC CTAAAAATGC CCTTTGCATG GACTTTTTTG ACTACCCTTT AGAAAATCAA
TTTGACACGA TTATTGGCAA TCCGCCCTAT GTCAAGCACA AGGATATTGC GCCAAGCACG
AAAGAAAAAC TCCATTACAG CCTTTTTGAT GAAAGGAGTA ATCTATACTT GTTTTTCATA
GAAAAAGCGA TCAAGCATTT AAAGCCTAAA GGCGAATTGA TTTTCATCAC CCCAAGGGAT
TTTTTAAAAT CCACTTCTAG CGTGAAATTA AACGAATGGA TTTACAAAGA AGGCACGATA
ACGCATTTTT TTGAATTAGG CGATCAAAAG ATTTTCCCAA ACGCCATGCC TAATTGCGTG
ATTTTTCGTT TTTGTAAAGG TGATTTCAGT AGAATCACCA ACGATGGTTT GCAATTTGTG
TGCAAAAAAG GCATTTTGTA TTTCCTCAAC CAATCTTACA CGCAAAAATT AAGCGAGGTT
TTTAAGGTTA AGGTGGGGGC AGTGAGCGGG TGCGATAAGA TTTTTAAAAA TGAAACATAC
GGGAATTTAG AATTTGTCAC CTCAATCACC AAAAGAACCA ATGTTTTAGA AAAAATGGTT
TTTGTCAATA AACCTAATGA TTATTTACTC CAGCATAAAG ACAGCTTGAT GCAAAGAAAG
ATTAAAAAAT TCAATGAAAG TAATTGGTTT GAATGGGGGA GGATGCATCA CATATCCCCT
AAAAAACGCA TTTATGTTAA CGCCAAAACG CGCCAAAAAA ACCCCTTTTT CATCCACCAA
TGCCCTAATT ATGACGGCTC TATTTTAGCG CTATTCCCTT ATAACCAAAA TTTGGATTTA
CAAAACCTCT GCGATAAACT CAACGCTATC AACTGGCAAG AATTAGGCTT TGTGTGCGGC
GGGCGTTTTT TGTTTTCGCA GCGCTCTTTA GAAAACGCCC TTTTGCCTAA AGACTTTTTA
AATTAG

FIG. 33A
SEQ ID NO:4

ATGGGTAAAT CTGAATTAAG TGGAAGATTA AATTGGCAAG CATTGGCTGG ATTAAAAGCT AGTGGTGCTG
AACAAAACTT ATATAACGTG TTTAACGCTG TTTTTGAAGG AACTAAATAC GTTTTATACG AGAAGCCAAA
GCACCTTAAA AATCTATACG CTCAAGTAGT CTTACCTGAT GATGTTATTA AAGAAATTTT TAATCCTTTA
ATTGATTTAT CAACTACTCA ATGGGGTGTT TCTCCAGATT TCGCAATAGA AAATACAGAA ACGCATAAAA
TTCTTTTTGG TGAAATTAAA AGACAAGATG GATGGGTAGA AGGTAAAGAT CCTAGTGCTG GCAGGGGTAA
TGCACATGAG AGATCTTGTA AATTATTTAC TCCTGGATTA TTAAAAGCTT ATAGAACAAT TGGTGGAATT
AACGATGAAG AGATATTGCC ATTCTGGGTT GTATTCGAAG GTGATATAAC ACGAGATCCC AAAAGAGTAA
GAGAAATTAC TTTCTGGTAT GACCACTATC AAGATAATTA TTTCATGTGG CGACCAAATG AATCAGGCGA
AAAATTAGTT CAACACTTCA ATGAAAAATT AAAAAAATAT TTAGATTAA

FIG. 33B
SEQ ID NO:5

MGKSELSGRL NWQALAGLKA SGAEQNLYNV FNAVFEGTKY VLYEKPKHLK NLYAQVVLPD DVIKEIFNPL
IDLSTTQWGV SPDFAIENTE THKILFGEIK RQDGWVEGKD PSAGRGNAHE RSCKLFTPGL LKAYRTIGGI
NDEEILPFWV VFEGDITRDP KRVREITFWY DHYQDNYFMW RPNESGEKLV QHFNEKLKKY LD

FIG. 34A

SEQ ID NO:6

ATGGCTATTA CATTATGTGA CATAAATGGT TGTAGACTTG AGAGAGGACA TACTGGTAAA CATAATAAAT TTCCTGAATT
TGTATGGACT TCTCAATTTA ATAAAAAAGA TATTGATAAG GTCAATAAAG CAGGATATGC AACACCAAGA GGTGGGGACA
AAGGAGCCTA TCAGAACCAT GTTTACAGAA ATAATAAAGT AATTATTCCT TTTGAAAGGT TGGAAAATGT TAATTTAAAT
AACTATCAAG ATGGATATGT TATTAGGTTA TTCCCTAATC AGTACTTTGA ATCAGCCGGG GTAGTTAAGC CGGAATTCTT
ACAACCAAAT TCATTTGTTA AAGTTGGGGA CAATGCATTT ATTTTATATC GCACACATTC ATCTTTTGAG GAATTACCTC
CTCTACCAGA CTGGGAGGTT AGACATCTAA AAAAGAACGG TAATATAGTT ACCAGAAGAA GTAAGGACGT AATCGATGCT
GGACATTATG TCTTACGATT ATCATCAATT AGTAACAAAA AAGAAAGAAA AGAGGGCCCT CCTCAAGGTA TTTTTGCACC
TGAATATGCA AATGCAGAGA CTAATTATCT GTCAAAAGCA TTTTTAGCCT GGTTAATTAT TAAAACTCAA AATAGTCCGT
ATAATGAAGA ACAATTCCAA CACTTAAGAG CGATCTTAAT TAGTCATAAT CTCATCAATA TTTCTCAACT TGAAGAAAAG
GCTATTCTAA AGAATGGTAT CACATGCTGC CCTTTATGCG AGCAAATTAT TTTTTACGAA CAGCTACACG AAATGGTTTC
TTTTGAAGGT GCGTCTGGCC TTGCGAATTC ACAAGAACAG GTTGAGGGTG CAACTAGGTC AACATCAGTT AATTTATTCC
ATATGGTACC ATTAGTATAT GAAACCTTGG AACACAAACC TGATCAAATA GCATGGGGCC ATGCCATTTG TAATACTAGA
CTTGGTCAAA GAGAGTGCCT GCCTCTTAGT AGACTAAAAC AAGAAGGTAC GCCCGTTGGT CTTCTTGATG AAGATTCGAA
TCTTGAAGTA TTAGGATGGA TTAGTAAAGA TAAGCAATTT ATTCGTACAG AAAATGGGGA AGTTTGGATT AAAATTACAG
ATATTGAATT TAACGATGAC TTTGAAGAAT AA

FIG. 34B

SEQ ID NO:7

MAITLCDING CRLERGHTGK HNKFPEFVWT SQFNKKDIDK VNKAGYATPR GGDKGAYQNH VYRNNKVIIP
FERLENVNLN NYQDGYVIRL FPNQYFESAG VVKPEFLQPN SFVKVGDNAF ILYRTHSSFE ELPPLPDWEV
RHLKKNGNIV TRRSKDVIDA GHYVLRLSSI SNKKERKEGP PQGIFAPEYA NAETNYLSKA FLAWLIIKTQ
NSPYNEEQFQ HLRAILISHN LINISQLEEK AILKNGITCC PLCEQIIFYE QLHEMVSFEG ASGLANSQEQ
VEGATRSTSV NLFHMVPLVY ETLEHKPDQI AWGHAICNTR LGQRECLPLS RLKQEGTPVG LLDEDSNLEV
LGWISKDKQF IRTENGEVWI KITDIEFNDD FEE

FIG. 35A

SEQ ID NO:8

ATGATTTTTG CTGATATTGA ATTTGAAAAA GAACTTTTTT CAGCTGCTAA TAAATTAAGG GGAAAAATTG CTCCAAGTGA
GTATAAGCAT TATGTTTTGC CTTTGATATT CCTTAGATAT TTATCTCTTA AATACCAACA AAGAAGGAAT GAAATTCAAC
AACAGATAAA TGATTCAAGG GATCACAAGA AAAATCAAGA TGAAGTGTTA AAGATATTGG AAGACAGGAC TGAATACACC
AAAGTAAATG TTTTCTATAT TCCTGAAAAA GCTAGTTGGG AATACTTATT GAAAAATTCC GAAAATGATA AAATTAAAGA
AATGATAGAT TCAGCTATGG AAATACTGGA AAATGAATAT GACGAGTTAA AAGGTGTTTT GCCAAAGATA TATAAAAACT
CAAATATACC GAATGAAGTT ATTAGTGATT TACTAAAACT ATTTTCTCAA GAAGTATTTT CAGCACATGA TGGAAGAAAT
GTTGATTTAT TGGGGAGAGT TTATGAATAC TTTATAAGTA ATTTTGCTAC TACAGAAGGT ACTAGAGGTG GTGAATATTT
TACACCGTCT TCAATCGTAA AATTATTGGT AGCAATGCTA GAGCCCATTA AAGGTACAGT TTATGATCCG GCCTGTGGGA
CAGGAGGAAT GTTTATTCAG TCTAATAAAT ATAGAGAAAA TAATCATAAC TTGTGTTTTG TAGGCCAGGA ACAAAACGAG
CTTACTATCA AATTGGCTAA AATGAATGGA ATTCTACATG GAATAAATCC TGAAATTAGA CAAGGTGATT CATTATTAAA
TGACCGTTAT CCAGAATTGA AAGCTGAAAT TGTAATATCT AATCCACCGT TTAATATGAA GGATTGGGGA GCTGAACGCC
TGCCACTTAA TGATAAGCGA TTAATAGGAC CGGTAACAAA CAGTAATGCA AATTACATGT GGATACAGCA TTTTCTATAC
CATTTAAAAG ATGGTGGTTT AGCAGGATTT GTTATTGCTA ATGGAGCTTT GACTAGTAAT CTGGCTGCTG AAAAAATTGT
AAGGAAACAC TTAATAGACA ATGATTATGT AGATTGTGTT GTTCAATTAC CTGAAAAAAT GTTCTTTGGT ACTGGCATTC
CAAGTGCTTT AGTGTTTTTA AGTAAGAATC GAAATGGAAG TAACGGCCAT GCCAAAAGAG AAAAAGAGGT TCTATTTATT
GATGCAAGCG ATAAGGGAAC ATTAGTGGGT AAAAAGAATA AAATATTTTT AGATGATGAA ATAAAAGAAA TTGCAGATTT
ATATCATTCA TTTAAATTTT TAAATGATAA TGATTATAAC CATAGTGGTT TTTACAAAAA GGTTAACATT GAAAAAATCG
TGGAAAATGA TTATAAATTA ACTCCAACTC TCTATGTAGG TGTAAAGGAA GAGACTGAAA TGGAGAAGCC ATTTAGAGAA
ATGATAATAG AATATAAAGC GATATTAGAG CAACAATTTG AAGAATCAAA CAAACTACAG CAGAAAATAT TAAAGAATTT
AGAGGGATTA TTATGA

FIG. 35B

SEQ ID NO:9

MIFADIEFEK ELFSAANKLR GKIAPSEYKH YVLPLIFLRY LSLKYQQRRN EIQQQINDSR DHKKNQDEVL KILEDRTEYT
KVNVFYIPEK ASWEYLLKNS ENDKIKEMID SAMEILENEY DELKGVLPKI YKNSNIPNEV ISDLLKLFSQ EVFSAHDGRN
VDLLGRVYEY FISNFATTEG TRGGEYFTPS SIVKLLVAML EPIKGTVYDP ACGTGGMFIQ SNKYRENNHN LCFVGQEQNE
LTIKLAKMNG ILHGINPEIR QGDSLLNDRY PELKAEIVIS NPPFNMKDWG AERLPLNDKR LIGPVTNSNA NYMWIQHFLY
HLKDGGLAGF VIANGALTSN LAAEKIVRKH LIDNDYVDCV VQLPEKMFFG TGIPSALVFL SKNRNGSNGH AKREKEVLFI
DASDKGTLVG KKNKIFLDDE IKEIADLYHS FKFLNDNDYN HSGFYKKVNI EKIVENDYKL TPTLYVGVKE ETEMEKPFRE
MIIEYKAILE QQFEESNKLQ QKILKNLEGL L

FIG. 36A

SEQ ID NO:10

ATGAAAAGTA CTTTGAAGGA ATATAAATTG GGTGATATTA CCGAAGTCGT TAATGGTGCC ACTCCTTCAA
CTAAAAAGCC TGAGTACTAT GAAAATGGTA CAATTCCATG GATTACTCCT AAAGATTTAT CAGGCTATTA
CTTTAAATAT ATATCTCATG GTGAACGTAA TATAACAGAG CTTGGTCTAA GAAATAGTTC AGCTAAGTTG
TTACCAAAAG GAACTGTATT ATTTTCCTCA AGAGCCCCAA TAGGATACGT AGCAATAGCT GATAATTGGT
TAACTACGAA CCAGGGATTT AAAAGTTTTA TATGTAATGA GGAGATTATT TACAATGAAT ACCTTTATTA
TTTTCTTATT GCTAAAAGGG ATTTTATTGA AACATTTGCG AATGGGAGTA CGTTTAAAGA GCTTTCATCA
ACTTCTGCAA AGAATATACC AATCAATCTT CCTAGTTTAG AAGAGCAAAA GAAGATTGTG ACAATTTTAG
GGGATTTGGA TAGAAAGATA GAATTAAATT ATAAAATTAT TGAAAGCTTA GAAAAAATAG CAGAAAGAAC
ATATAAATAT TGGTTTGTCG ATGAATTAAA TCAAGATGAA CAGCACATCC GTAATGGATG GGAAACTGCT
AAAATTGGCG ATGTGGTGGA ACTTTTGGGA GGGGGAACCC CTAAAACTTC GGAAAGTAAG TATTGGGAAG
ATGGAGATAT TAATTGGTTT ACTCCTTCAG ATTTAACAAA AACTAGACAG CTTTTTGTAC GTGATTCTCA
AAGAAAAATA ACAATTGATG GACTTAATAA CAGTGCAGCG AAATTAATTC CCCCTTATTC CTTGTTAATG
TCAAGTAGAG CTACAATTGG CGAGTTGGCA ATTAATCAAG AATCTGCTAC TACAAATCAA GGGTTTATTG
TATTAATACC AAATGAAAAA ATTTCTATTT ACCAATTATA CTTTTGGGCT AAACTTAATA AGAGCAAAAT
TATTTCAATG GCAAATGGTA GTACTTTTAA AGAAATTAGT AAGCGGGATT TTAAATCTTT GGAGATAATA
TTACCAAAAA ATATAGACAC TTTTAATTCA ATTATGCAAG ATTATTTTAG GAAAATTGAG GAGTTAATTG
ATGAAATAAA AATCTTAAAA ACCGCAAGAG ATAATTTAAT TCCAAAACTT ATAAAATGA

FIG. 36B

SEQ ID NO:11

MKSTLKEYKL GDITEVVNGA TPSTKKPEYY ENGTIPWITP KDLSGYYFKY ISHGERNITE
LGLRNSSAKL LPKGTVLFSS RAPIGYVAIA DNWLTTNQGF KSFICNEEII YNEYLYYFLI
AKRDFIETFA NGSTFKELSS TSAKNIPINL PSLEEQKKIV TILGDLDRKI ELNYKIIESL
EKIAERTYKY WFVDELNQDE QHIRNGWETA KIGDVVELLG GGTPKTSESK YWEDGDINWF
TPSDLTKTRQ LFVRDSQRKI TIDGLNNSAA KLIPPYSLLM SSRATIGELA INQESATTNQ
GFIVLIPNEK ISIYQLYFWA KLNKSKIISM ANGSTFKEIS KRDFKSLEII LPKNIDTFNS
IMQDYFRKIE ELIDEIKILK TARDNLIPKL IK

FIG. 37A

SEQ ID NO:12

ATGAAACAGT TTGCAGATCC TTTTGAAAGA AGATTCCTTG ATGCAATTGA ACATCATCTT GATGGAATTT
CTGAGAAAAT AAAAAAAGAC TTTACACACA AAAACTTTTT AAAAGAATTG AATGGCCTTA AAGGTGATAA
AGTCTATCAT GACTTAGGCT TTGATACCGC TGAATATACT CTGGTACGTC TTATAGGAAG AATGAGCATA
AGCGTTGGGA GAAGGCTGGG GGAGATATAC GATAAAGTCC CTCGTTATGT TGCTGCCGCG CGATTTGGTC
TTCAACCAAA TCAAATTGCA GAAGTATTTG ATGGTCTTGA GTTAGATATA GCTTTGCGCA ATAGCCTTTT
GTCAGATGAT GATAAAATTC ACATAAAAAA AATAACTGAA AAGATGTCAG GCGAAACATA CTCGGGAATC
GGAATCGAAA TTCGTTATAA CTTTAATCCA AATGACAGTT CCCGTTTAAG AAAAGACGTC GATGTAGCTT
CTAAATTGTC GGCCGCGGGG TTATTTCCTG TTTATTTAAT ATTTAGCTCT CTCAGTCCTA GGAATGATGC
AATAGCCCGT CTTAAAAGAG GGGGATGGAG CTTTAAACAG GGGCAGGAAG CCTTAGACTT CCTTACCGAA
CTTTTAGGAG TGGATATTGG GTCTGTTTTA TCTGACCCAA TAATAGCCGC AGAAACTAGG GAGAAAACAT
CAAAAATTAT GAAGTCTATA TTTGAATCAG AGGCATTCCA ATCTGTTATA CCGGGAGAGT GGAGTAAACT
ATAG

FIG. 37B

SEQ ID NO:13

MKQFADPFER RFLDAIEHHL DGISEKIKKD FTHKNFLKEL NGLKGDKVYH DLGFDTAEYT LVRLIGRMSI
SVGRRLGEIY DKVPRYVAAA RFGLQPNQIA EVFDGLELDI ALRNSLLSDD DKIHIKKITE KMSGETYSGI
GIEIRYNFNP NDSSRLRKDV DVASKLSAAG LFPVYLIFSS LSPRNDAIAR LKRGGWSFKQ GQEALDFLTE
LLGVDIGSVL SDPIIAAETR EKTSKIMKSI FESEAFQSVI PGEWSKL

FIG. 38A

SEQ ID NO:14

ATGACAAATT TTTCGCACTC AGCTCTAACG AGCTACGATC TTCTCGGGCA TGAAATTGTC CAAGATTCTG
AAGCTGTTAG CTCGGGTCCA TATCTGGTCA GCTATGACCC GATCCCTGTA CGTCGGTCTA CATTCCTAGC
TGGACTGTCA GAGAACGTTC ACTCGTGGTT TCGTCTCACA CCAAGTTTCG GACCGGATCT AGTTCGAACA
ATCATCAAAC AGATGAATCT TGCGCCGCAC TCACACATCC ATGACCCTTT CTCAGGAGCC GGGACTACCG
CGATTGAGGC TTCGTTAGAG GGCTATGAAG CAAGCTGCGT AGAAGTTAAT CCGTTTCTCT ACTTCGTGGG
GAAAACATCC ATAGATTGGT CTATCAATGC TGATGATGCT GCAGCGCAGC TAGAAAGCAT TAAAAATAAA
TATTATAGCA TGTCTGCAAC CGCTACTTTG GATAACATAG CCGACCTAGG AATAGATATA CCAAAAATAC
ACAATATTCA TCGGTGGTGG AGAAACGATG TTCTTAAAGA TATATTAGTC CTAAAATCTT CTATCAGATC
TTGCACACAA GATAAGTATT GTTCCTTTTT TGAGCTAGCC CTAGCTGCAG TTCTCGTTCC AGATTTGACA
AATGTAACGC TAGGAAAACT ACAACTGCAC TTTGTAAACA AAGACGATAA AGAGATAAAC GTCTGGCCTA
CATATGAATC TCATGCAAAA AAAATGATTC ACGACTTGTC ATTAATTAAT AAGCAAAATT TCGAATTTTT
GCCCAAGATT ATTTATGGTG ATTCAACTCA AAAATCAACA TTTAGCGAGG TGGCAGGGAT AGATGCTATA
ATAACATCCC CTCCGTACCC TAATAGGTAC AGCTATATTT GGAATACTCG CCCTCACCTG TACATTCTTG
ATATGATTTC CGAAGCAAAA GAGGCTTCGC AAATAGATCG TAGAACGATT GGTGGAACAT GGGGGACAGC
AACTTCCGAA TTAGGAAAGG GTATATTTTC TCCAATCAAT GCTGTAGTCA AAGACGCGCT TGAAGGGGTT
CACGAAAGAA TCGCCGGTTC CGATCAACTC ATGGCAAACT ATGTAACTCA TTATTTTAAT CGGCTCTTTT
TACATATAGA AGCTATAAAA CCATCACTTA ATCCAAAAGC AAAGCTTGCT TATGTTGTTG GGAACTCTTG
GATTAAGGGC GAATATGTAG CCACTGACGT AATCTTAGCA AAAATTATCG AAGGGGCTTT GCCAGGCTCA
TCAATTGATG GTCTTCATCG TTTCCGTCGC CGGAACAGTG GAAAGAATCT CTTTGAAACT ATAGTTTACT
CCACTCTCCC GGTATAA

FIG. 38B

SEQ ID NO:15

MTNFSHSALT SYDLLGHEIV QDSEAVSSGP YLVSYDPIPV RRSTFLAGLS ENVHSWFRLT PSFGPDLVRT
IIKQMNLAPH SHIHDPFSGA GTTAIEASLE GYEASCVEVN PFLYFVGKTS IDWSINADDA AAQLESIKNK
YYSMSATATL DNIADLGIDI PKIHNIHRWW RNDVLKDILV LKSSIRSCTQ DKYCSFFELA LAAVLVPDLT
NVTLGKLQLH FVNKDDKEIN VWPTYESHAK KMIHDLSLIN KQNFEFLPKI IYGDSTQKST FSEVAGIDAI
ITSPPYPNRY SYIWNTRPHL YILDMISEAK EASQIDRRTI GGTWGTATSE LGKGIFSPIN AVVKDALEGV
HERIAGSDQL MANYVTHYFN RLFLHIEAIK PSLNPKAKLA YVVGNSWIKG EYVATDVILA KIIEGALPGS
SIDGLHRFRR RNSGKNLFET IVYSTLPV

FIG. 39

SEQ ID NO:17

>M1.EarI CTCTTC 1245 nt

GTGAATCAGA AAAATGAAAA ATCATTTATG CGTTTGCAAT CAACCTTTAG CGGTGGCAAA
GGTAGTCCAA TGCATGATTG GTACCCATAT TTAGAGGGTT ATTCTCCCGA ATTTGTGAAA
TGCTTGATTT CACGATTTGC TCCTAAAGCC AAAACAATTT TAGATCCATT TTGTGGCTCT
GGAACAACAG CCATTGTTTC CGTTTTAGAG GGTTTAAATA ATTACTATTG CGAAGTAAAC
CCTTTATGCC AATATATTAT TGAAACTAAA CTAATAGCTT TAACATTAAG CGAAGAAGAA
AAAACAAAAT TAGTAAATGA ACTTTATTCT ATTTCTAATG AAATAACTAA TGTACTCAAA
CCTTCTGCAA CCGAGACAGA TCTAGAGAAA TCATTTAAAT CCGTTTTTGG TAATACGAAA
TTTTTTGAGG ATCACATATT TAAAGATATA CTTAGTTATC AATGTTACAT TAGCTCTATC
GAAGATGAAA ATCTTAAGAG ACTTCTGACA ATAGCAGGGA TTAGATCGTT AATCCCTTCC
TCGTTATTGG TAAGACGAGG TGATTTACGA TTCAAGACAC AAAAAGAATT AGAGAAAGGC
AACCAGGGCT TTCGCTTTCA TGTACAAAAA AGCTTAGAAT TAATTGCCAG TGATTTATTA
GACATTACGG AAGGTAGTGG TTTAGCTACC TTCTTATGTG ATGATGCCAA AGAAATATCT
GGGAATAACC TGATTGATGC TGTAATAACA AGCCCGCCAT ATTTAAATGG CACAAATTAT
TTTAGAAATA CTAAAATTGA ACTTTGGTTT ATAGGGAAAT TAAAGACCAA ATCAGATCTA
AGACATTATA GGGATTTAGC TATTACCAGT GGTATTAACG ATGTAACTAA AGGTAAAAGC
TTATCTTCAA ATAATACTAT TATCTCAGAA ATACCATTAT TATCTGAATG TATTAAAGAA
CTAAGCATAA AAGAGTATGA TAGTCGTATT TCAATGATGG TTGAAAACTA CTTTTGGGAC
ATGTTCAAAT TCTTATCAAA ACTCCCAAAA TTACTAACTA ATGATGCGAC TATCTGTATA
GATTTAGGTG ATTCTGTTTA TTGTAACGTC TACATCCCTA CACAAGATAT TTTGAAAGAA
ATGATGTCAA AGTTAGGTTT TGAAGAGAAC GAAAGGGTCA TTCTTCGTGA ACGAAAATCC
CGCAATGGAA CAAAGTTAGT CCAGACTGTT CAGGTTTTTA AATGA

FIG. 40

SEQ ID NO:18

>M2.EarI CTCTTC 1140 nt

ATGAAAAATA AATATTTTAG TAAAAAATGG GAGCAATTCA AGAAAGAATT ACCCCATCAA
TCAGGTGAAA TGGTAAAGAG AAATTGGGGC CATAACTGGC ACTCTATGTG TTCATACCAA
GGGAAACTTA AACCATCAAT AGCTAGATCT TTAATTGATA CATTCATGCC ATCAAGTAAG
GGACGTATAT TAGATGTCTT CTCAGGTGTT GGCACCATTC CTTTCGAAGC AAGATTACTT
GGTCATACTG CATATGGATT TGATATTAGT CCAGCAGCAG TTAATATTTC ACGCGCAAAA
CTAGAAGTTA TAAGTAAAAA TGAAATCCAA GAGGTAATTA ATAAATTATC TGATTTTATT
GAGCAAAACA AAAATTCAAT AGATTATAAC GAACATAATT TAATAAGGTT TAATGGTTCA
ATTGAATCCT ATTTTCATCC TGAAACTTTT AAGGAAATAC TGTGTGCTCG TAAATTCTTT
TTAATAAAAG GTGAATTAAA TGCATCTGAA TCGTTAGTAC AGTCATGTTT ATTACATATT
TTACATGGTA ATCGTCCGTA TGCATTGAGT AGAAAGTCCC ATCCTATTAC ACCTTTCGCG
CCTACTGGAG ATTTTATATA CAGTAATTTA GTTATAAAGT TAATCAAAAA AGTTGAAAGA
GTCTTGCAAA ATTCTGATGG TATCCCAGAT ACTGGCAGCA AAGTATTTTA TCAGGACTCT
ACAAAAAGTT GGCCTGAAGA AGTAAATAAT TTAGATGCAA TTATAACATC ACCTCCATTT
TATGATAGTA CCCGTTTCTA TTCAGCAAAT TGGATGCGAT TATGGTTTTC TGGTTGGGAA
AAAGATGACT TCCAAACGAA GCCAAAAGAT TTTGTGGACG AAACTCAGAA AAAAAGCTTT
GAAATATATG ATAATATATT CAAACAATCT CAACAATGCT TAAAAAAAGA TGGCGTTTTT
TTAATGCACG TTGGCAAAAG TAAAAAAAGT GATATGGCAG GACAAATTGC TAAAATTGGT
AGTAATTATC TTAGCCTTAT AGATATATTT GACGAAAGTG TTGAACATTG CGAAAGTCAC
GGAATTAAAG ACAAAGGCAC GACAACCCAT CATCAGTACC TTGTCTTTAC GAAAGATTAG

FIG. 41A

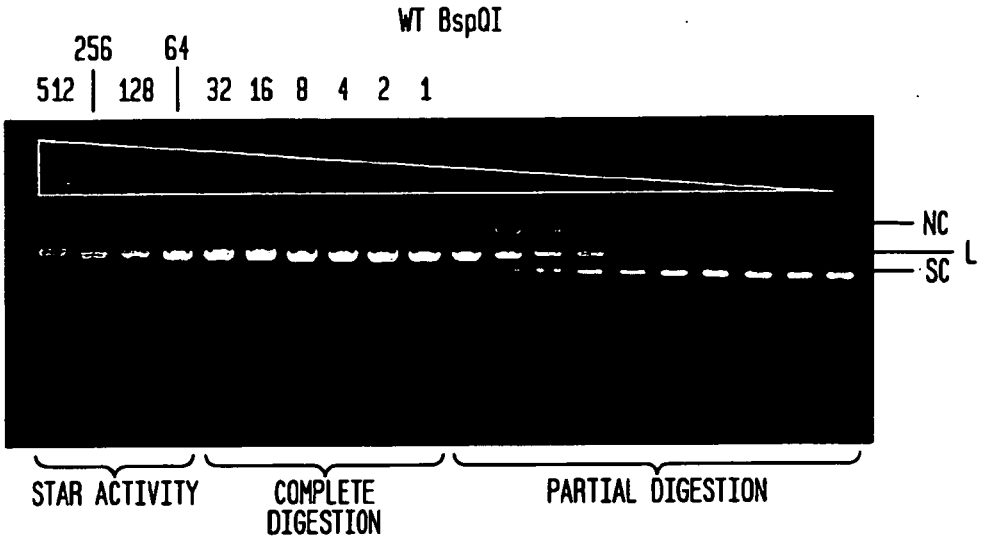


FIG. 41B

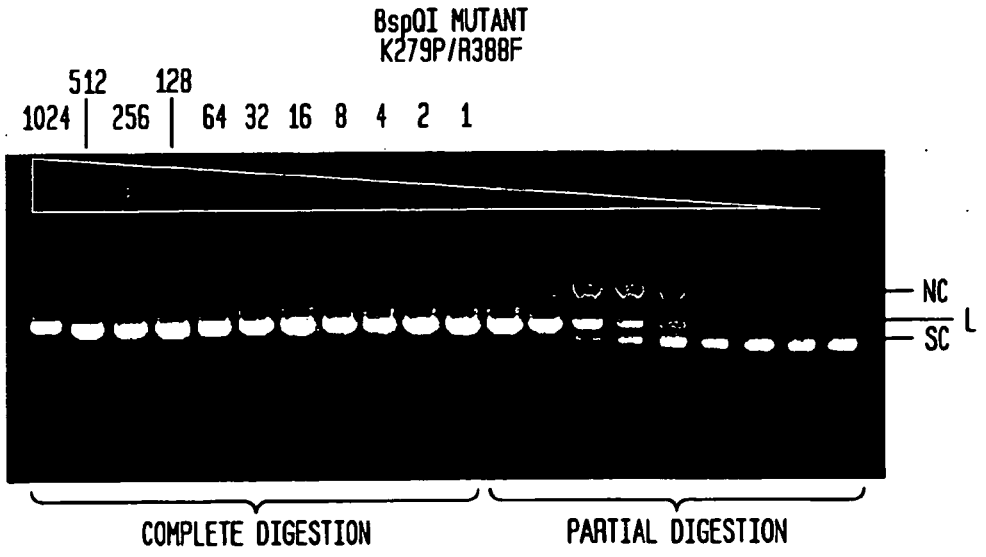


FIG. 42

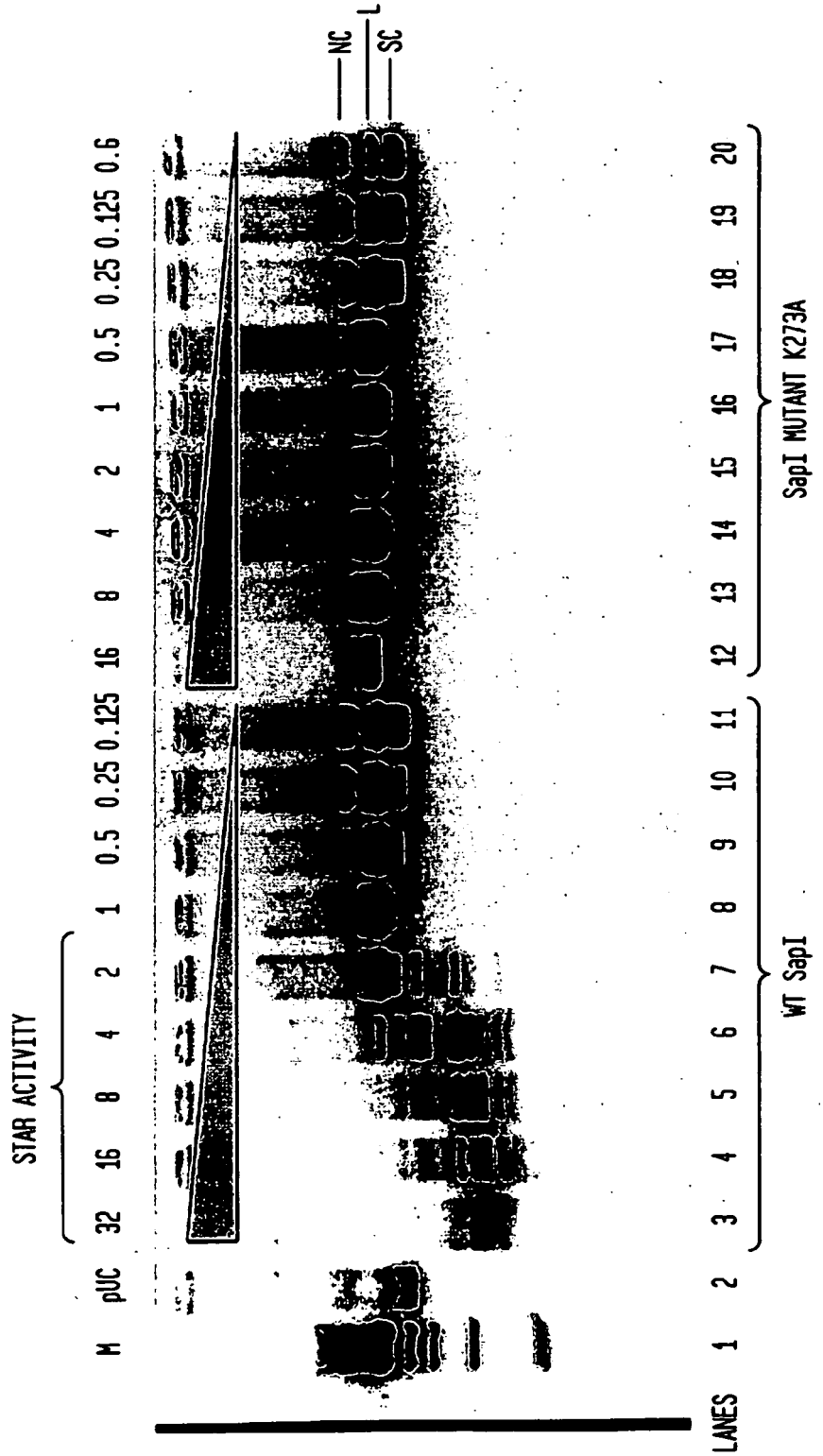
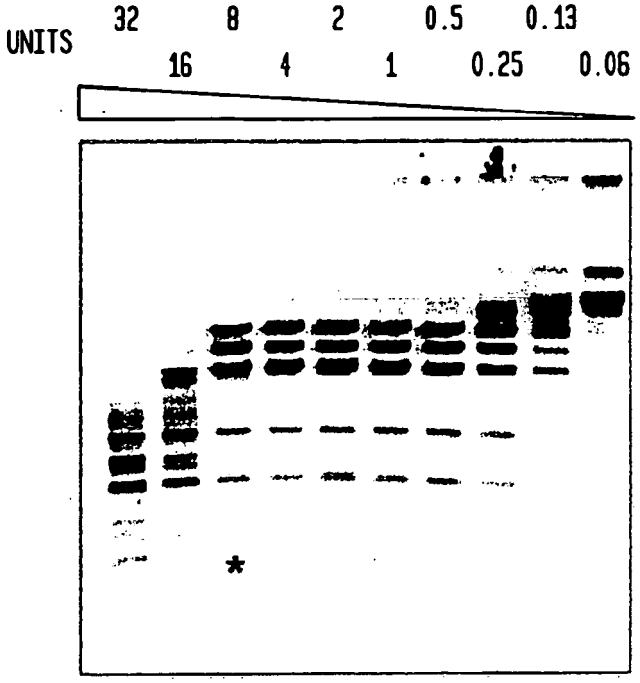
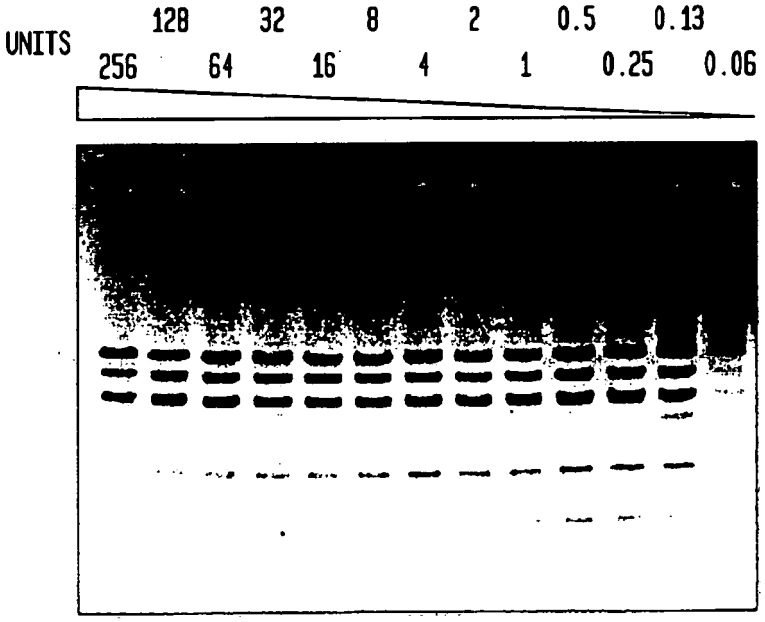

NC
L
SC
LANES 1 2 3 4 5 6 7 8 9 10 11 12 13 14 15 16 17 18 19 20
M pUC 32 16 8 4 2 1 0.5 0.25 0.125 16 8 4 2 1 0.5 0.25 0.125 0.6
STAR ACTIVITY
WT SapI
SapI MUTANT K273A

FIG. 43A



WT KpnI

FIG. 43B



KpnI D16N/E132A/D148E

FIG. 44A

>AgeI ACCGGT 272 aa (SEQ ID NO:79)

MRLDLDFGRG LVAHVMLDNV SEEQYQQISD YFVPLVNKPK LKSRDAIGQA
FVMATEVCPD ANPSDLWHHV LYRIYIREKI GTDPSQSWVR TSGEAFEVAL
VERYNPVLAR HGIRLTALFK GQKGLALTRM GVADRVGSRK VDVMIEKQGG
GRSPDAEGFG VVGGIHAKVS LAERVSDDIP ASRIMMGEGL LSVLSTLDVK
SFPPPHGDLV NRGELGTPDR PSDKANYIEG HGDFSACFSY NLRTPPSNAT
TPSGRHIYVS ASLVRTTSSP TT

>AvrII CCTAGG 358 aa (SEQ ID NO:80)

MEEDLDLSEN IEAASAELTT LYQVAADAMK DYIEIYLALS KQSDGFSNIN
NLDLTSRNRR LVVIHGLSLE LDPDTSTPEE IKREAERMLA IALDTESAIT
AGVYEKMRLF ASSLVDQLFE QTDELNSLSS EYLSANPGFL PFFQQLAGLR
SKSELKREVG NASDNSISKA VAERILERII RNLRIRTFSK EKLLQAVEPT
LEGIVRDLVG KVLLENIVAD ALSDLQVPFM RESEYQSLKG VIYDFRADFV
IPDAQNPIAF IEVRKSSTRH ASLYAKDKMF SAINWKGKNK RLLGILVVEG
PWTRETLRVM ANVFDYVTPL TRVSQVAEAI RAYLDGDKTR LKWLVNFSIE
EADHDNIT

>AvrII CCTAGG 1077 nt (SEQ ID NO:110)

ATGGAAGAAG ACCTTGATTT ATCTGAAAAT ATCGAAGCTG CATCTGCGGA
GCTTACGACT CTTTATCAGG TAGCTGCTGA TGCTATGAAA GATTATATTG
AAATCTATCT TGCGCTGAGT AAACAGTCTG ATGGGTTTTC AAATATTAAC
AATCTTGACT TAACTTCTCG TAACAGGCGT TTGGTAGTTA TACATGGACT
TTCGTTAGAG TTAGATCCAG ATACTTCGAC TCCAGAGGAA ATTAAACGTG
AAGCTGAACG AATGCTAGCG ATAGCTCTTG ATACAGAGTC AGCAATTACG
GCAGGAGTAT ATGAAAAAAT GCGTCTCTTC GCAAGCTCTT TAGTAGATCA
GCTATTTGAA CAAACGGATG AACTTAATTC ATTATCATCG GAATATTTGT
CAGCAAATCC AGGATTTTTG CCGTTTTTCC AGCAGTTGGC GGGGCTTAGA
AGTAAATCAG AGTTAAAGAG AGAAGTAGGA AATGCCTCTG ACAATAGTAT
TTCTAAAGCG GTTGCAGAGA GAATATTAGA GCGCATTATA CGTAACTTGA
GAATTCGCAC TTTTTCCAAA GAGAAACTAT TACAAGCTGT TGAGCCTACT
TTAGAAGGAA TAGTCAGGGA TCTCGTAGGA AAAGTGTTAT TGGAAAATAT
AGTTGCTGAT GCTTTATCTG ATTTACAAGT TCCTTTCATG CGTGAATCAG
AGTATCAAAG CCTTAAAGGA GTGATTTATG ATTTCCGCGC TGATTTTGTG
ATACCAGACG CACAAAATCC AATTGCTTTT ATCGAGGTGC GAAAAAGCTC
TACACGACAT GCGTCACTCT ATGCCAAGGA TAAGATGTTT TCAGCGATTA
ATTGGAAAGG AAAAAATAAA AGGCTTTTGG GTATTTTGGT TGTGGAAGGA
CCTTGGACAA GAGAAACTCT TCGCGTCATG GCAAATGTGT TTGATTACGT
TACACCTTTA ACTCGTGTTT CCCAAGTTGC AGAAGCTATC AGAGCATATC
TAGATGGGGA TAAAACGAGA CTGAAGTGGT TAGTTAATTT CAGTATTGAA
GAAGCAGACC ACGACAACAT AACCTAA

FIG. 44B

>BsmBI CGTCTC 530 aa (SEQ ID NO:81)

MAKYGRGKFL PHQNYIDYMH FIVNHKNYSG MPNAIGEDGR INWQVSSGKT
TSFYEYYQAR FEWWEKKADE LNLPGTGNSN KAFSLAARLI HPTGQRPCRL
CGKYQYVGYM YVSHNLYKRW SKITGREDLF FKKQNIIEAA NIFKSIMGEQ
ALINELTTIF PERKDYFNRL PNIEDFFVSS SHIKNNGNYI SPGFMANPPD
RLDGFHDYGI CCRKEKDPGR HDDNMRLYNH DRRAFMWWSE GDWALADALY
NKAGAGKCAD PDCQKEVEKI SPDHVGPISC GFKQIPFFKP LCASCNSAKN
RRFSYQDVKE LLKYENYTGD SVASWQVRAL WDNCKHLVKN DDDSKLLSNL
MRSLQDYYLR SLYKLFSNGF AHLLSYFLTP EYAHYKITFE GLNTSTLEYE
RYYKTFKKTK STSSLAARIV RIAFEELEIY NSKDINERKL IKFDTSSWEK
DFENIISYAT KNLSLDEEAS KWNKVLTDKN LSSTEKDKKI SSLLEDKNYE
VYKKQFYILK DLLVEHFNKI GEQIAKDYMK

>BspQI GCTCTTC 430 aa (SEQ ID NO:16)

1 MRRLAKNSRN DSYLSNRDYQ EIVRENTTTI SFPLKEKHTL TLTKKIGLNQ 50
51 TAGFGGWFFP DSPCLLTVTV LSSFGTKVTS KTFSLSKDWN RVGLAWINEH 100
101 SSDTMSIVLE FSDVEIVHTW GLTCDVFNVH ELIIDAIEDQ NKLIDVLNQE 150
151 HLSPETYYLN HDSDTDLIEN LESTEEIKIV NQSQKQISLK KCCYCQRYMP 200
201 VNILVRSNSS FHKHKSKKTG FQNECRACKK WRINNSFNPV RTKDQLHESA 250
251 VITREKKILL KEPEILQKIK NRNNGEGLKS IIWKKFDKKC FNCEKELTIE 300
301 EVRLDHTRPL AYLWPIDEHA TCLCEKCNNT KHDMFPIDFY QGDEDKLRRL 350
351 ARITGLDYES LVKRDVNEVE LARIINNIED FATNVEARTF RSIRNKVKEV 400
401 RPDTDLFEIL KSKNINLYNE LQYELLTRKD 430

>BspQI GCTCTTC 1293 nt (SEQ ID NO:101)

ATGAGACGAT TAGCAAAAAA TTCACGGAAC GACAGTTATT TAAGTAATAG
GGATTACCAG GAAATCGTGA GGGAAAATAC CACTACAATA TCGTTTCCCT
TAAAAGAAAA ACATACTCTG ACTTTAACGA AAAAAATAGG GCTAAATCAG
ACTGCTGGAT TCGGAGGATG GTTTTTCCCT GATTCACCAT GTTTATTAAC
AGTAACTGTA CTATCCTCTT TCGGTACAAA GGTAACTTCT AAAACCTTTA
GCCTTTCTAA AGATTGGAAT CGTGTTGGGC TTGCTTGGAT TAACGAGCAT
TCGAGTGACA CCATAAGCAT TGTCCTAGAG TTTAGTGATG TGGAAATAGT
TCATACATGG GGACTTACAT GTGATGTTTT TAATGTCCAT GAATTAATTA
TTGATGCTAT AGAAGATCAA AATAAACTAA TAGACGTGCT AAATCAAGAA
CATTTATCTC CTGAAACATA TTATTTAAAC CATGACTCTG ATACTGATTT
AATTGAGAAT TTGGAATCTA CAGAAGAGAT AAAGATAGTT AACCAAAGCC
AAAAGCAAAT CTCTTTAAAA AAATGCTGTT ATTGTCAACG TTATATGCCT
GTGAACATAT TAGTTCGTTC AAATTCATCA TTTCATAAAC ACAAGAGTAA
GAAAACTGGT TTTCAAAATG AATGTCGGGC TTGTAAGAAG TGGAGAATAA
ATAATTCATT CAATCCAGTC AGAACAAAAG ACCAACTACA TGAATCAGCA
GTTATTACAC GTGAAAAAAA AATATTACTT AAAGAACCTG AAATATTACA
GAAAATCAAA AATAGAAATA ACGGTGAGGG CTTAAAAAGT ATTATATGGA

FIG. 44C

AAAAATTTGA TAAAAAATGC TTTAATTGTG AAAAAGAATT AACCATTGAA
GAGGTACGCC TAGACCATAC AAGACCACTT GCTTATCTGT GGCCTATCGA
TGAACACGCA ACTTGTTTAT GTGAAAAATG CAACAATACA AAACATGATA
TGTTTCCTAT CGATTTTTAT CAAGGGGACG AAGACAAATT AAGACGTTTA
GCTAGAATTA CGGGGTTAGA TTATGAATCT CTAGTTAAGA GGGACGTAAA
TGAAGTTGAA CTTGCAAGAA TAATCAATAA CATTGAAGAC TTTGCAACTA
ATGTAGAGGC ACGTACTTTT CGCTCAATAA GAAATAAAGT AAAAGAAGTA
CGTCCCGATA CTGACCTATT TGAAATTCTT AAATCTAAAA ATATTAATTT
ATATAATGAA CTTCAATATG AACTTCTTAC CCGTAAGGAT TAA

>EagI CGGCCG 301 aa (SEQ ID NO:82)

MKKRRDLVEV FGYNPMDLSP EVRALWNLGA CPFLNKECIK INHDQTIIYG
TCSVTSPYGD VIICPNRLYA NDYETLHKVS RDAFGDDVPF LTYSNFIKYR
ATYKDCIVAL GKNSGKEVQV GRALSMDWVL VRITDGELKE YVGVEIQSID
ITGNYRDAWH AYKNLKPIDI IDNLPTSQHG LNWANVHKRL IPQIIRKGVV
YSRSNYVKKG LYFILPEIVY NKFEDVIGAD IPLLKTQTNK SITVHTYSLG
EPAANGEQRK LISEREIIFD LDEFSKRFTT GPNLPKGDDL DAVIKKALGM
M

>EcoRI GAATTC 277 aa (SEQ ID NO:83)

MSNKKQSNRL TEQHKLSQGV IGIFGDYAKA HDLAVGEVSK LVKKALSNEY
PQLSFRYRDS IKKTEINEAL KKIDPDLGGT LFVSNSSIKP DGGIVEVKDD
YGEWRVVLVA EAKHQGKDII NIRNGLLVGK RGDQDLMAAG NAIERSHKNI
SEIANFMLSE SHFPYVLFLE GSNFLTENIS ITRPDGRVVN LEYNSGILNR
LDRLTAANYG MPINSNLCIN KFVNHKDKSI MLQAASIYTQ GDGREWDSKI
MFEIMFDIST TSLRVLGRDL FEQLTSK

>EcoRV GATATC 245 aa (SEQ ID NO:84)

MSLRSDLINA LYDENQKYDV CGIISAEGKI YPLGSDTKVL STIFELFSRP
IINKIAEKHG YIVEEPKQQN HYPDFTLYKP SEPNKKIAID IKTTYTNKEN
EKIKFTLGGY TSFIRNNTKN IVYPFDQYIA HWIIGYVYTR VATRKSSLKT
YNINELNEIP KPYKGVKVFL QDKWVIAGDL AGSGNTTNIG SIHAHYKDFV
EGKGIFDSED EFLDYWRNYE RTSQLRNDKY NNISEYRNWI YRGRK

>HindIII AAGCTT 300 aa (SEQ ID NO:85)

MKKSALEKLL SLIENLTNQE FKQATNSLIS FIYKLNRNEV IELVRSIGIL
PEAIKPSSTQ EKLFSKAGDI VLAKAFQLLN LNSKPLEQRG NAGDVIALSK
EFNYGLVADA KSFRLSRTAK NQKDFKVKAL SEWREDKDYA VLTAPFFQYP
TTKSQIFKQS LDENVLLFSW EHLAILLQLQ LEETNIFPFE QLWNFPKKQS
KKTSVSDAEN NFNRDFNKYF MDLFKIDKDT LNQLLQKEIN FIEERSLIEK
EYWKKQINII KNFTREEAIE ALLKDINMSS KIETIDSFIK GIKSNDRLYL

FIG. 44D

>HpaI GTTAAC 254 aa (SEQ ID NO:86)

MKYEEINFKV PVESPYYPNY SQCVIERIYS ILRNQKDMGD DRIIINTNLK
KGLPLENINK IAGPMIEAWA EEVFSGIRDN RDNQYNLINV EAQERLGISD
IILQFQVNNN VITGNVDVKA TSNDIPDSGK SPNITSFSRI RTAYVKDPNF
IFIILSIKHS VYVKRNEYTN LMDGIMQIID FNVYDLKYIS DSDISYNPAL
GTGQIQIKDI HYVSSQKRTT WQMCQLLDLK YLRSKKRTIE QFYNEAKRNK
WIKD

>KpnI GGTACC 218 aa (SEQ ID NO:87)

MDVFDKVYSD DNNSYDQKTV SQRIEALFLN NLGKVVTRQQ IIRAATDPKT
GKQPENWHQR LSELRTDKGY TILSWRDMKV LAPQEYIMPH ATRRPKAAKR
VLPTKETWEQ VLDRANYSCE WQEDGQHCGL VEGDIDPIGG GTVKLTPDHM
TPHSIDPATD VNDPKMWQAL CGRHQVMKKN YWDSNNGKIN VIGILQSVNE
KQKNDALEFL LNYYGLKR

>NcoI CCATGG 288 aa (SEQ ID NO:88)

MATAPGHLLG QIIGNVMEEA LKPVLQEMAD RHDLYLDSKG LRPGVRSGAL
VTWTDDLGNN HDLDFVLERG GSATKAGNPA AFIEAAWRRY TKHSKAKAQE
IQGAVLPVLA AWNNVKPTPA AVVAGQWTAP SLQQMRSNGF VVLHLHFPTT
AQVFGGNGIN IEGTGEGTPD AFWQQQCDAY TSKSEADKDS LATALRTAHA
QEFRTFVAEL ERRVVRAIDY VVVTPLHGHG SQYTSIENAI EAVRTYSCGE
ESAPFLRFEI RISYTNGDVI QATFGSSSDA IEFLDTFN

>NheI GCTAGC 328 aa (SEQ ID NO:89)

MSSYHDDLNI LNVDFNHLRL TELIKLADQA EPFYLWVEKI FRQVSGRADS
LETIIEVEER VVLKMAILTC FTSDEKELPK LFNGVGVPYP HIKACYFFFA
WLVRDAATQR LDPLIREAFT QLKSIHPQMK KTELESEIFS QLLVNYRNEL
IHFSWPVIRE VLISRLEGSR RAARGSYLEL FVRTALAQSI TYFYKIYGNY
GKFLDVKIHD KPLKVKNRTY DVVAELIGNN HNTQYLILPV KTRETQGGGH
AHLFTRDIEQ SNNDIRELYP NAVIAPVIIA ENWSDTEKDL ENVGYNDIFH
FSVNPNRFAG FSDVEQIRLN RLVERILL

FIG. 44E

>NotI GCGGCCGC 383 aa (SEQ ID NO:90)

MRSDTSVEPE GANFIAEFFG HRVYPEVVST EAARNDQATG TCPFLTAAKL
VETSCVKAET SRGVCVVNTA VDNERYDWLV CPNRALDPLF MSAASRKLFG
YGPTEPLQFI AAPTLADQAV RDGIREWLDR GVHVVAYFQE KLGGELSISK
TDSSPEFSFD WTLAEVESIY PVPKIKRYGV LEIQTMDFHG SYKHAVGAID
IALVEGIDFH GWLPTPAGRA ALSKKMEGPN LSNVFKRTFY QMAYKFALSG
HQRCAGTGFA IPQSVWKSWL RHLANPTLID NGDGTFSLGD TRNDSENAWI
FVFELDPDTD ASPRPLAPHL EIRVNVDTLI DLALRESPRA ALGPSGPVAT
FTDKVEARML RFWPKTRRRR STTPGGQRGL FDA

>PstI CTGCAG 326 aa (SEQ ID NO:91)

MKELKLKEAK EILKALGLPP QQYNDRSGWV LLALANIKPE DSWKEAKAPL
LPTVSIMEFI RTEYGKDYKP NSRETIRRQT LHQFEQARIV DRNRDLPSRA
TNSKDNNYSL NQVIIDILHN YPNGNWKELI QQFLTHVPSL QELYERALAR
DRIPIKLLDG TQISLSPGEH NQLHADIVHE FCPRFVGDMG KILYIGDTAS
SRNEGGKLMV LDSEYLKKLG VPPMSHDKLP DVVVYDEKRK WLFLIEAVTS
HGPISPKRWL ELEAALSSCT VGKVYVTAFP TRTEFRKNAA NIAWETEVWI
ADNPDHMVHF NGDRFLGPHD KKPELS

>PvuII CAGCTG 157 aa (SEQ ID NO:92)

MSHPDLNKLL ELWPHIQEYQ DLALKHGIND IFQDNGGKLL QVLLITGLTV
LPGREGNDAV DNAGQEYELK SINIDLTKGF STHHHMNPVI IAKYRQVPWI
FAIYRGIAIE AIYRLEPKDL EFYYDKWERK WYSDGHKDIN NPKIPVKYVM
EHGTKIY

FIG. 44F

>SacI GAGCTC 358 aa (SEQ ID NO:93)

MGITIKKSTA EQVLRKAYEA AASDDVFLED WIFLATSLRE VDAPRTYTAA
LVTALLARAC DDRVDPRSIK EKYDDRAFSL RTLCHGVVVP MSVELGFDLG
ATGREPINNQ PFFRYDQYSE IVRVQTKARP YLDRVSSALA RVDEEDYSTE
ESFRALVAVL AVCISVANKK QRVAVGSAIV EASLIAETQS FVVSGHDVPR
KLQACVAAGL DMVYSEVVSR RINDPSRDFP GDVQVILDGD PLLTVEVRGK
SVSWEGLEQF VSSATYAGFR RVALMVDAAS HVSLMSADDL TSALERKYEC
IVKVNESVSS FLRDVFVWSP RDVHSILSAF PEAMYRRMIE IEVREPELDR
WAEIFPET

>SalI GTCGAC 315 aa (SEQ ID NO:94)

MINADKPHRW NDDVQASVRL YNQWFLDAAP KAYRDTRQLT IDEVEQAFQR
TANMTSITPE VLKAHPKTLA TLRMSTAPPI ARDRLVGLSH GSKSLLDTME
KGKLPPRMKG DVLDTHLAKM CAVLTDLLQL DLFHWYPTGE PAEPRQRELA
ATVVADRLCG AIADPIVRNA QERRQLALIE EWLLARGYTK KTHSASLPLN
TMQPGTFSFR QNVVVGSDLP VNIPVDAVIQ PHTPHSHKLP ILIEAKSAGD
FTNTNKRRKE EATKIHQLQL KYGNEISLTL FLCGYFNTGY LGYSAAEGLD
WVWEHRIDDL EAAGA

>SapI GCTCTTC 432 aa (SEQ ID NO:95)

MRRLATQRRE DAYKSNRDYQ TVHEAQSLRV NSTDDDNLSL FLLKDISPRE
DSKNIVGFGG FVKPEIATTM ALTLTTDIDK QIKSVPLSSN WNRISIVAKF
ASNPSVSITL GFDQTPWVDF WGINSDDIGL SFVSDAVPLE MSMIDSIHIA
PETLYLDHSS ACLLDIDPVE STRFKTGHGD PLSLKKCSYC GRLLPIDLER
PGKLSFHKHR AKITNHQNEC RSCKKWRINN SFNPMRTIDQ LNESALITRE
RKIFLQEPEI LQEIKDRTGA GLKSQVWERF HRKCFNCRKD LKLSEVQLDH
TRPLAYLWPI DEHATCLCAQ CNNTKKDRFP VDFYSEQQIR ELSDICGLPY
QDLCARSLNL DQLDRIERNI AEFSKEWDVR TFASTARRIS EVYPARDLFE
TLKKESESAY NKIIEKLKER PDALLDEALP LD

>SbfI CCTGCAGG 323 aa (SEQ ID NO:96)

MNSSDGIDGT VASIDTARAL LKRFGFDAQR YNVRSAVTLL ALAGLKPGDR
WVDSTTPRLG VQKIMDWSGE HWAKPYATGS REDFRKKTLR QWVDNGFAVL
NADNLNIATN SQLNEYCLSD EALQALRAYG TEGFEESLVV FLDEASKAVK
ARAEALQAAM ISVDLPGGEE FLLSPAGQNP LLKKMVEEFV PRFAPRSTVL
YLGDTRGKHS LFEREIFEEV LGLTFDPHGR MPDLILHDEV RGWLFLMEAV
KSKGPFDEER HRSLQELFVT PSAGLIFVNC FENRESMRQW LPELAWETEA
WVAEDPDHLI HLNGSRFLGP YER

FIG. 44G

>ScaI AGTACT 227 aa (SEQ ID NO:97)

MINDQLPRWV REARVGTRTG GPAMRPKTSD SPYFGWDSED WPEVTRQLLS
EQPLSGDTLV DAVLASWESI FESRLGSGFH IGTQIRPTPQ IMGFLLHALI
PLELANGDPS WRADLNSSEK DLVYQPDHKY SIEMKTSSHK DQIFGNRSFG
VENPGKGKKA KDGYYVAVNF EKWSDAPGRL PRIRTIRYGW LDHTDWVAQK
SQTGQQSSLP AVVSNTQLLA IHTGGQR

>SphI GCATGC 235 aa (SEQ ID NO:98)

MTSKDPIVLS ADQIAWLRQL KMSKRAALVR DYILEYGAVT TGKLAELGYS
HPPRAARDLK DAGAGVVTIM VKGPDGRRMA SYAFNGKANE DGAGRVVIPK
AFGEALKRAH GGKCAVCYGD FSERELQCDH RVPFAIAGDK PKLVQEDFMP
LCASDNRAKS WSCENCPNWE LKDEDTCRSC FWASPENYTH VSTRPERRIN
LLFQGDEVEI FDALKNAAAN EGVSLTEATK RKLAD

>SspI AATATT 281 aa (SEQ ID NO:99)

MSKAAYQDFT KRFSLLIKKH PNLITMTLSN IFTMRLIGNK THGDLAEIAI
SEFINQYMYD FKSIHVGKDL YRAKSKEEDI TVENEITKEK FPISLKAYGD
GPLQLSTDKN FLMYPLLEEI GAFINAKEKI EEIFANEAFS CFSEINVLPL
IYDEKRQRCN ILVFDAARAR AETAYIRKET EGSGRKHPAY RFFDKNKNYI
CEVRYGNAAA NALQRGLWTN TKNATSFFDS VTNGWVDYSH NLVLVKLLSH
ALVSSRKGHE AALEEIKKDI LQLKQTNGIN V

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Application Number
EP 11 07 5102

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Category	Citation of document with indication, where appropriate, of relevant passages	Relevant to claim	CLASSIFICATION OF THE APPLICATION (IPC)
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TECHNICAL FIELDS SEARCHED (IPC)

C12N

The present search report has been drawn up for all claims

Place of search	Date of completion of the search	Examiner
Munich	19 October 2011	Keller, Yves

CATEGORY OF CITED DOCUMENTS

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